

12

EUROPEAN PATENT APPLICATION

21 Application number: 88810622.6

22 Date of filing: 13.09.88

51 Int. Cl.⁴: **C 07 D 401/14**
C 07 D 401/12,
C 07 D 211/94, C 09 D 7/12,
C 08 K 5/34, C 08 K 5/35

30 Priority: 21.09.87 US 99418 21.09.87 US 99412
21.09.87 US 99413

43 Date of publication of application:
29.03.89 Bulletin 89/13

84 Designated Contracting States:
BE DE FR GB IT NL

71 Applicant: CIBA-GEIGY AG
Klybeckstrasse 141
CH-4002 Basel (CH)

72 Inventor: Winter, Roland A.E.
23 Banksville Road
Armonk New York 10504 (US)

Galbo, James P.
119 East Hartsdale Avenue
Hartsdale New York 10530 (US)

Mar, Andrew
97 Beacon Street
Norwalk Connecticut 06851 (US)

Behrens, Rudolf A.
P.O. Box 8213
New Fairfield Connecticut 06812 (US)

Malherbe, Roger François, Dr.
Bleichstrasse 7
CH-4058 Basle (CH)

54 Stabilization of coatings with N-hydroxy hindered amines.

57 N-hydroxy-substituted hindered amine light stabilizers impart outstanding stabilization properties to coatings based on a variety of resins, the stabilized coatings exhibiting improved durability, weatherability, and the like. Some of them are new compounds.

EP 0 309 401 A1

Description

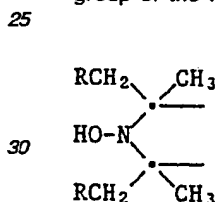
STABILIZATION OF COATINGS WITH N-HYDROXY HINDERED AMINES

The instant invention relates to the stabilization of a wide variety of ambient curable or acid catalized thermosetting coating composition by the incorporation therein of N-hydroxy-substituted hindered amine light stabilizers. The invention further relates to the new compounds of this type of hindered amines.

Hindered amine light stabilizers are well known to be effective in stabilizing a host of organic substrates including polymers against the deleterious effects of light and oxygen. Such hindered amine light stabilizers have been used in the stabilization of hot-crosslinkable alkyd or acrylic metallic stoving lacquers (US 4,426,472) and in stabilizing acid-catalyzed stoving lacquers based on hot-crosslinkable acrylic, polyester or alkyd resins (US 4,344,876 and 4,426,471). The hindered amine light stabilizers of these patents do not possess structures having a hydroxyl group substituted directly on the hindered N-atom of the compound.

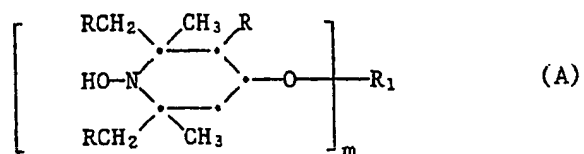
Related hindered amine stabilizers have been utilized individually and in combination with ultra-violet light absorbers to improve the performance characteristics of coating systems. Notwithstanding such improvements, there still exists a need to further retard the photooxidation and photodegradation of coating systems and thereby provide increased effectiveness by maintaining the physical integrity of the coatings. Such effectiveness can be manifested by prevention of embrittlement, cracking, corrosion, erosion, loss of gloss, chalking and yellowing of the coating.

It has now been determined that the aforementioned improvements can be achieved by substitution on the hindered N-atom of the hindered amines with hydroxyl groups and the utilization of such derivatives in ambient curable and acid catalyzed thermosetting coating systems. In particular, the physical integrity of the coatings is maintained to a higher degree with significant reduction in loss of gloss and yellowing. Accordingly, the present invention relates to a stabilized ambient curable or acid catalyzed thermosetting coating composition containing an effective stabilizing amount of a hydroxyl substituted hindered amine compound containing the group of the formula

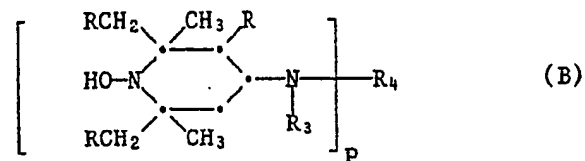


where R is hydrogen or methyl.

More particularly, the instant invention to the use of a hindered amine derivative corresponding to one of formulae A to M

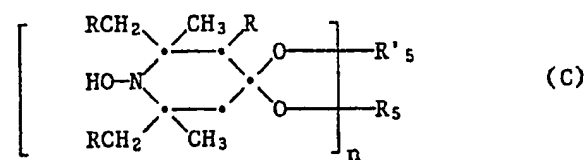


5

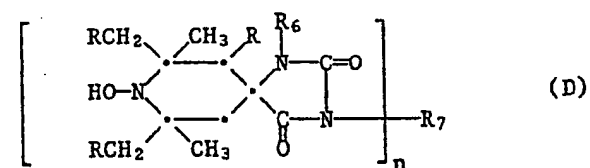


10

15

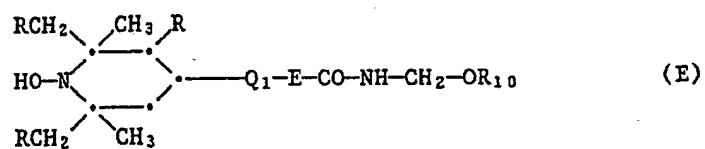


20



25

30



35

40

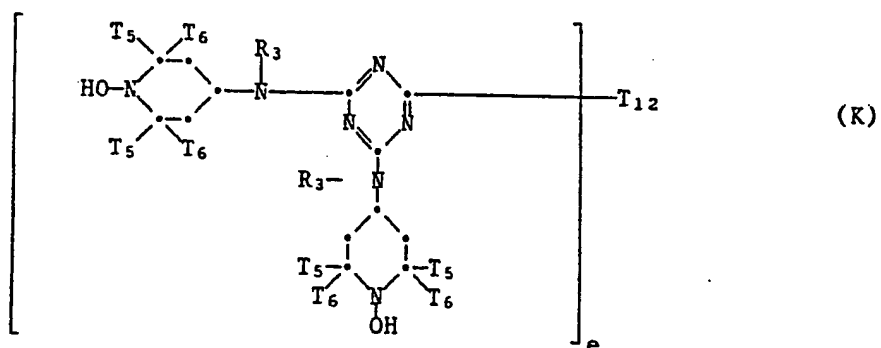
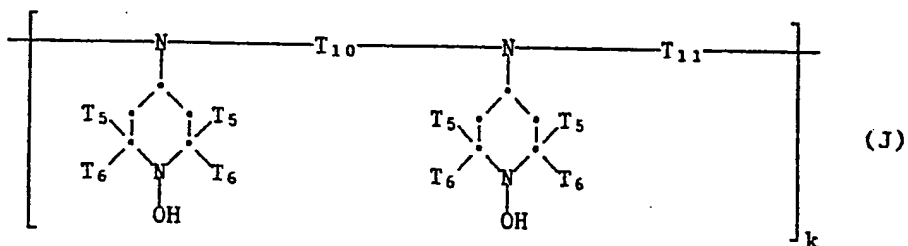
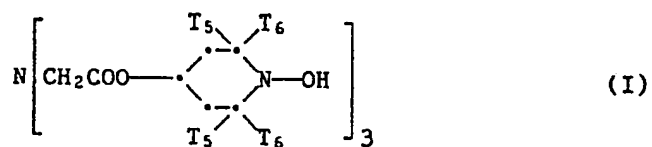
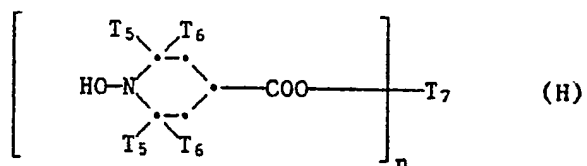
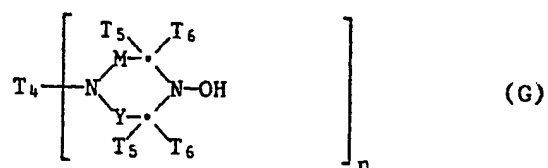
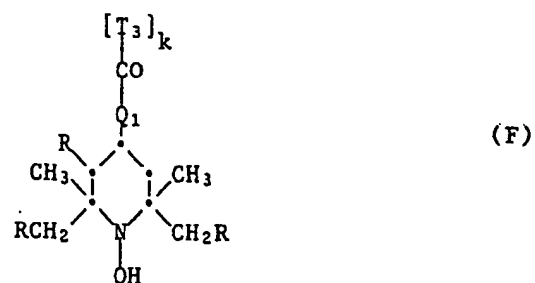
45

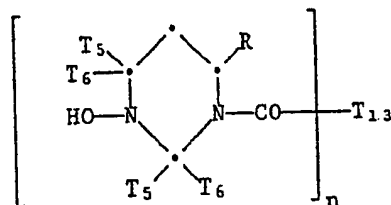
50

55

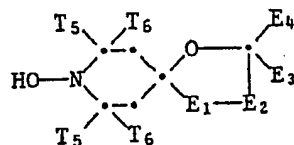
60

65





(L)



(M)

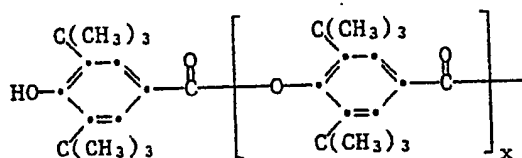
wherein

R is hydrogen or methyl,

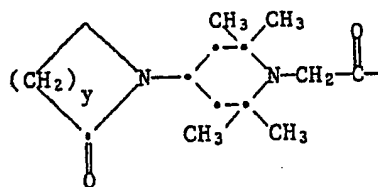
m is 1-4,

when m is 1,

R₁ is hydrogen, C₁-C₁₈ alkyl optionally interrupted by one or more oxygen atoms, C₂-C₁₂ alkenyl, C₆-C₁₀ aryl, C₇-C₁₂ aralkyl, glycidyl, a monovalent acyl radical of an aliphatic, cycloaliphatic, araliphatic or aromatic carboxylic or carbamic acid, preferably an acyl radical of an aliphatic carboxylic acid having 2-18 C atoms, or a cycloaliphatic carboxylic acid having 5-12 C atoms or an aromatic carboxylic acid having 7-12 C atoms; or R₁ is a group



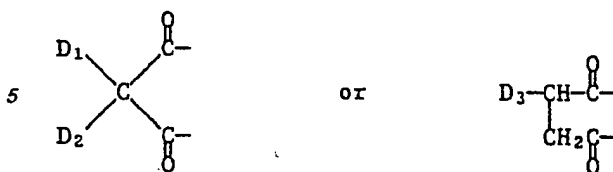
wherein x is 0 or 1, or is a group



wherein y is 2-4;

when m is 2,

R₁ is C₁-C₁₂ alkylene, C₄-C₁₂ alkenylene, xylylene, a divalent acyl radical of an aliphatic, cycloaliphatic, araliphatic or aromatic dicarboxylic or dicarbamic acid, preferably an acyl radical of an aliphatic dicarboxylic acid having 2-18 C atoms, of a cycloaliphatic or aromatic dicarboxylic acid having 8-14 C atoms, or of an aliphatic, cycloaliphatic or aromatic dicarbamic acid having 8-14 C atoms; or R₁ is a group



10

wherein D₁ and D₂ are independently hydrogen, an alkyl radical containing up to 8 carbon atoms, phenyl, benzyl or 3,5-di-*t*-butyl-4-hydroxybenzyl and D₃ is an alkyl or alkenyl radical containing up to 18 carbon atoms; when m is 3, R₁ is a triavalent acyl radical of an aliphatic, unsaturated aliphatic, cycloaliphatic, or aromatic tricarboxylic acid;

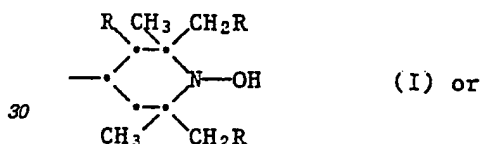
15 when m is 4, R₁ is a tetravalent acyl radical of a saturated or unsaturated aliphatic or aromatic tetracarboxylic acid including 1,2,3,4-butanetetracarboxylic acid, 1,2,3,4-but-2-enetetracarboxylic acid, and 1,2,3,5- and 1,2,4,5-pentanetetracarboxylic acid;

p is 1, 2 oder 3,

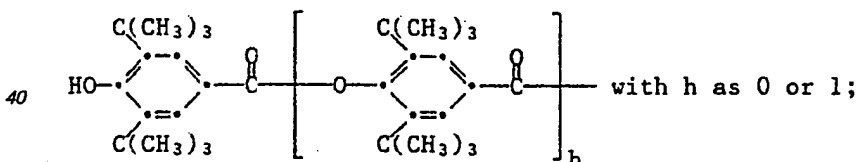
20 R₃ is hydrogen, C₁-C₁₂ alkyl, C₅-C₇ cycloalkyl, C₇-C₉ aralkyl, C₂-C₁₈ alkanoyl, C₃-C₅ alkenoyl or benzoyl; when p is 1,

R₄ is hydrogen, C₁-C₁₈ alkyl, C₅-C₇ cycloalkyl, C₂-C₈ alkenyl unsubstituted or substituted by cyano, carbonyl or carbamide group, aryl, aralkyl, or it is glycidyl, a group of the formula -CH₂-CH(OH)-Z or -CONH-Z wherein Z is hydrogen, methyl or phenyl; or R₄ is a group of formula I

25



35

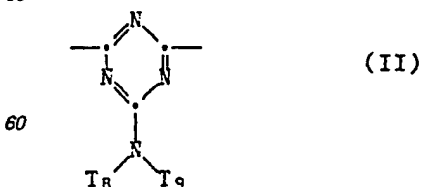


45

or R₃ and R₄ together are alkylene of 4 to 6 carbon atoms or 2-oxoalkylene or the cyclic acyl radical of an aliphatic or aromatic 1,2- or 1,3-dicarboxylic acid, when p is 2,

50 R₄ is C₂-C₁₂ alkylene, C₆-C₁₂ arylene, xylylene, a -CH₂CH(OH)-CH₂- group, or a group -CH₂-CH(OH)-CH₂-O-X-O-CH₂-CH(OH)-CH₂- wherein X is C₂-C₁₀ alkylene, C₆-C₁₅ arylene or C₆-C₁₂ cycloalkylene; or, provided that R₃ is not alkanoyl, alkenoyl or benzoyl, R₄ can also be a divalent acyl radical of an aliphatic, cycloaliphatic or aromatic dicarboxylic acid or dicarbamic acid, or can be the group -CO-, or R₄ is a group of formula II

55



65 where T₈ and T₉ are independently hydrogen, alkyl of 1 to 18 carbon atoms or a group of formula I, or T₈ and T₉

together are alkylene of 4 to 6 carbon atoms or 3-oxapentamethylene, preferably T₈ and T₉ together are 3-oxapentamethylene;

when p is 3,

R₄ is 2,4,6-triazinetriyl,

n is 1 or 2 and

when n is 1,

R₅ and R'₅ are independently C₁-C₁₂ alkyl, C₂-C₁₂ alkenyl, C₇-C₁₂ aralkyl, or R₅ is also hydrogen, or R₅ and R'₅ together are C₂-C₈ alkylene or hydroxyalkylene or C₄-C₂₂ acyloxyalkylene;

when n is 2,

R₅ and R'₅ together are (-CH₂)₂C(CH₂)₂;

R₆ is hydrogen, C₁-C₁₂ alkyl, allyl, benzyl, glycidyl or C₂-C₆ alkoxyalkyl;

when n is 1,

R₇ is hydrogen, C₁-C₁₂ alkyl, C₃-C₅ alkenyl, C₇-C₉ aralkyl, C₅-C₇ cycloalkyl, C₂-C₄ hydroxyalkyl, C₂-C₆ alkoxyalkyl, C₆-C₁₀ aryl, glycidyl, a group of the formula -(CH₂)_t-COO-Q or of the formula -(CH₂)_t-O-CO-Q wherein t is 1 or 2, and Q is C₁-C₄ alkyl or phenyl; or

when n is 2,

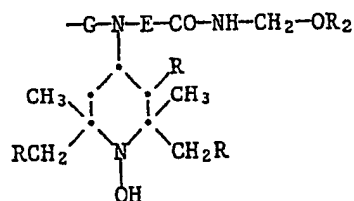
R₇ is C₂-C₁₂ alkylene, C₆-C₁₂ arylene, a group -CH₂CH(OH)-CH₂-O-X-O-CH₂-CH(OH)-CH₂- wherein X is C₂-C₁₀ alkylene, C₆-C₁₅ arylene or C₆-C₁₂ cycloalkylene, or a group -CH₂CH(OZ')CH₂-(OCH₂-CH(OZ'))CH₂- wherein Z' is hydrogen, C₁-C₁₈ alkyl, allyl, benzyl, C₂-C₁₂ alkanoyl or benzoyl;

Q₁ is -N(R₈)- or -O-;

E is C₁-C₃ alkylene, the group -CH₂-CH(R₉)-O- wherein R₉ is hydrogen, methyl or phenyl, the group -(CH₂)₃-NH- or a direct bond;

R₁₀ is hydrogen or C₁-C₁₈ alkyl,

R₈ is hydrogen, C₁-C₁₈ alkyl, C₅-C₇ cycloalkyl, C₇-C₁₂ aralkyl, cyanoethyl, C₆-C₁₀ aryl, the group -CH₂-CH(R₉)-OH wherein R₉ has the meaning defined above, or R₈ is a group of the formula I or a group of the formula



wherein G is C₂-C₆ alkylene or C₆-C₁₂ arylene, or R₈ is a group -E-CO-NH-CH₂-OR₁₀;

Formula F denotes a recurring structural unit of a polymer where T₃ is ethylene or 1,2-propylene, or is the repeating structural unit derived from an alpha-olefin copolymer with an alkyl acrylate or methacrylate; preferably a copolymer of ethylene and ethyl acrylate, and where k is 2 to 100;

T₄ has the same meaning as R₄ when p is 1 or 2,

T₅ is methyl,

T₆ is methyl or ethyl, or T₅ and T₆ together are tetramethylene or pentamethylene or a mixture of said derivatives, preferably T₅ and T₆ are each methyl,

M and Y are independently methylene or carbonyl, preferably M is methylene and Y is carbonyl, and T₄ is ethylene where n is 2;

T₇ is the same as R₇, and T₇ is preferably octamethylene when n is 2,

T₁₀ and T₁₁ are independently alkylene of 2 to 12 carbon atoms, or T₁₁ is a group of formula II;

e is 2, 3 or 4 and

T₁₂ is a group -N(R₅)-(CH₂)_d-N(R₅)-

or $-\text{NH}(\text{CH}_2)_a-\text{N}(\text{CH}_2)_b-\text{N}[(\text{CH}_2)_c-\text{N}]_f\text{H}$

where a, b and c are independently 2 or 3, d is 2 to 10 and f is 0 or 1, preferably a and c are each 3, b is 2 and f is 1;

T₁₃ is the same as R₄ with the proviso that T₁₃ cannot be hydrogen when n is 1;

E₁ and E₂, being different, each are -CO- or N(E₅)-, preferably E₁ is -CO- and E₂ is -N(E₅)- where E₅ is hydrogen, C₁-C₁₂ alkyl or alkoxycarbonylalkyl of 4 to 22 carbon atoms;

E₃ is hydrogen, alkyl of 1 to 30 carbon atoms, phenyl, naphthyl, said phenyl or said naphthyl substituted by chlorine or by alkyl of 1 to 4 carbon atoms, or E₃ is phenylalkyl of 7 to 12 carbon atoms, or said phenylalkyl substituted by alkyl of 1 to 4 carbon atoms, and

E₄ is hydrogen, alkyl of 1 to 30 carbon atoms, phenyl, naphthyl or phenylalkyl of 7 to 12 carbon atoms, or

E₃ and E₄ together are polymethylene of 4 to 17 carbon atoms, or said polymethylene substituted by up to four alkyl groups of 1 to 4 carbon atoms, preferably methyl.

Of particular interest are coating compositions containing a hindered amine derivative of formula A, B, G-K or M wherein R is hydrogen and T₅ and T₆ are methyl.

5 In the structures A to M, if any substituents are C₁-C₁₈ alkyl, they are for example methyl, ethyl, n-propyl, n-butyl, sec-butyl, tert-butyl, n-hexyl, n-octyl, 2-ethylhexyl, n-nonyl, n-decyl, n-undecyl, n-dodecyl, n-tridecyl, n-tetradecyl, n-hexadecyl or n-octadecyl.

If R₁ is a monovalent acyl radical of a carboxylic acid, it is for example an acyl radical of acetic acid, stearic acid, salicylic acid, methacrylic acid, benzoic acid or β-(3,5-di-tert-butyl-4-hydroxyphenyl) propionic acid.

10 If R₁ is a divalent acyl radical of a dicarboxylic acid, it is for example an acyl radical of adipic acid, succinic acid, suberic acid, sebacic acid, phthalic acid, maleic acid, butyl-(3,5-di-tert-butyl-4-hydroxybenzyl)-malonic acid, or bicycloheptenedicarboxylic acid.

If R₁ is a divalent acyl radical of a dicarbamic acid, it is for example an acyl radical of hexamethylenedicarbamic acid or of 2,4-toluylenedicarbamic acid.

15 If any substituents are C₅-C₇ cycloalkyl, they are in particular cyclohexyl.

As C₇-C₉ aralkyl, R₃ is particularly phenethyl or above all benzyl.

As C₂-C₁₈ alkanoyl, R₃ is for example propionyl, butyryl, octanoyl, dodecanoyl, hexadecanoyl, octadecanoyl, but preferably acetyl; and as C₃-C₅ alkenoyl, R₃ is in particular acryloyl.

If R₄ is C₂-C₆ alkenyl unsubstituted or substituted by a cyano, carbonyl or carbamide group, it is for example 20 1-propenyl, allyl, methallyl, 2-butenyl, 2-pentenyl, 2-hexenyl, 2-octenyl, 2,2-dicyanovinyl, 1-methyl-2-cyano-2-methoxycarbonyl-vinyl or 2,2-diacetylaminovinyl.

If any substituents are C₂-C₁₂ alkylene, they are for example ethylene, propylene, 2,2-dimethylpropylene, tetramethylene, hexamethylene, octamethylene, decamethylene or dodecamethylene.

If any substituents are C₆-C₁₅ arylene, they are for example o-, m- or p-phenylene, 1,4-naphthylene or 25 4,4'-diphenylene.

As C₆-C₁₂ cycloalkylene, X is especially cyclohexylene.

If R₅ is C₂-C₈ alkylene or hydroxyalkylene, it is for example ethylene, 1-methyl-ethylene, propylene, 2-ethylpropylene or 2-ethyl-2-hydroxymethylpropylene.

As C₄-C₂₂ acyloxyalkylene, R₅ is for example 2-ethyl-2-acetoxymethylpropylene.

30 If any substituents are C₂-C₆ alkoxyalkyl, they are for example methoxymethyl, ethoxymethyl, propoxymethyl, tert-butoxyethyl, ethoxyethyl, ethoxypropyl, n-butoxyethyl, tert-butoxyethyl, isopropoxyethyl or propoxypropyl.

If R₇ is C₃-C₅ alkenyl, it is for example 1-propenyl, allyl, methallyl, 2-butenyl or 2-pentenyl.

As C₇-C₉ aralkyl, R₇ is in particular phenethyl or above all benzyl; and as C₅-C₇ cycloalkyl, R₇ is especially 35 cyclohexyl.

If R₇ is C₂-C₄ hydroxyalkyl, it is for example 2-hydroxyethyl, 2-hydroxypropyl, 3-hydroxypropyl, 2-hydroxybutyl or 4-hydroxybutyl.

As C₆-C₁₀ aryl, R₇ is in particular phenyl, or alpha- or β-naphthyl which is unsubstituted or substituted by halogen or C₁-C₄ alkyl.

40 If R₇ is C₂-C₁₂ alkylene, it is for example ethylene, propylene, 2,2-dimethylpropylene, tetramethylene, hexamethylene, octamethylene, decamethylene or dodecamethylene.

If R₇ is C₆-C₁₂ arylene, it is for example o-, m- or p-phenylene, 1,4-naphthylene or 4,4'-diphenylene.

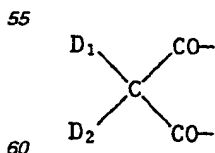
If Z' is C₂-C₁₂ alkanoyl, it is for example propionyl, butyryl, octanoyl, dodecanoyl or preferably acetyl.

As C₅-C₇ cycloalkyl, R₈ is in particular cyclohexyl.

45 As C₆-C₁₀ aryl, R₈ is particularly phenyl, or alpha- or β-naphthyl which is unsubstituted or substituted with halogen or C₁-C₄ alkyl. As C₁-C₃ alkylene, E is for example methylene, ethylene or propylene.

As C₂-C₆ alkylene, G is for example ethylene, propylene, 2,2-dimethylpropylene, tetramethylene or hexamethylene; and as C₆-C₁₂ arylene, G is o-, m- or p-phenylene, 1,4-naphthylene or 4,4'-diphenylene.

Preferred are coating compositions containing a hindered amine derivative of formula A wherein R is 50 hydrogen, m is 1, 2 or 4 and when m is 1, R₁ is an acyl radical of an aliphatic carboxylic acid having 12-18 C atoms or of an aromatic carboxylic acid having 7-15 C atoms and when m is 2, R₁ is a divalent acyl radical of an aliphatic dicarboxylic acid having 4-18 C atoms or of an aromatic dicarboxylic acid having 8-14 C atoms or R₂ is a group



wherein D₁ is C₁-C₈ alkyl or 3,4-di-tert-butyl-4-hydroxybenzyl and D₂ is D₁ or hydrogen, and when m is 4, R₁ is 65 tetravalent acyl radical of a butane- or pentanetetracarboxylic acid.

The following compounds are examples of hindered amine derivatives applicable for use in the invention:

1. 1,4-dihydroxy-2,2,6,6-tetramethylpiperidine
2. 1-hydroxy-4-salicyloxy-2,2,6,6-tetramethylpiperidine
3. 1-hydroxy-4-methacryloyloxy-2,2,6,6-tetramethylpiperidine
4. 1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl 3,5-di-tert-butyl-4-hydroxyhydrocinnamate
5. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) sebacate 5
6. di-(1-hydroxy-2,3,6-trimethyl-2,6-diethyl-piperidin-4-yl) phthalate
7. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl n-butyl-(3,5-di-tert-butyl-4-hydroxybenzyl)malonate
8. hexane-1',6'-bis-(4-carbamoyloxy-1-hydroxy-2,2,6,6-tetramethylpiperidine)
9. N,N'-bis(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)hexamethylene-1,6-diacetamide
10. 4-(N-cyclohexylacetamido)-1-hydroxy-2,2,6,6-tetramethylpiperidine 10
11. 4-methacrylamido-1-hydroxy-2,2,6,6-tetramethylpiperidine
12. N-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)-epsilon-caprolactam
13. N-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)-succinimide
14. N-(1-hydroxy-2,3,6-trimethyl-2,6-diethyl-piperidin-4-yl)-maleimide
15. 8-aza-2,7,9,9-pentamethyl-8-hydroxy-1,4-dioxaspiro[4.5]decane 15
16. 9-aza-3-hydroxymethyl-3-ethyl-9-hydroxy-8,8,10,10-tetramethyl-1,5-dioxaspiro[5.5]undecane
17. 3-n-octyl-1,3,8-triaza-8-hydroxy-7,7,9,9-tetramethyl-spiro[4.5]decane-2,4-dione
18. 8-hydroxy-2,7,7,9,9-pentamethyl-2-hexyl-1-oxa-3,8-diazaspiro[4.5]decane-4-one
19. 3-hydroxy-2,2,4,4-tetramethyl-7-oxa-3,20-diazaspiro[5.1.11.2]heneicosan-21-one
20. 1,1'-ethylenebis-(4-hydroxy-3,3,5,5-tetramethylpiperazin-2-one) 20
21. 1,1'-sebacoyl-bis-(3-hydroxy-2,2,4,4,6-pentamethylhexahydropyrimidine)
22. N-hydroxy derivative of polycondensation product of 2,4-dichloro-6-tert-octylamino-s-triazine and 4,4'-hexamethylenebis-(amino-2,2,6,6-tetramethylpiperidine)
23. N,N',N''-tetrakis[4,6-bis(butyl-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)amino)-s-triazin-2-yl]1,10-diamino-4,7-diazadecane 25
24. N-hydroxy derivative of polycondensation product of 2,4-dichloro-6-morpholino-s-triazine and 4,4'-hexamethylenebis-(amino-2,2,6,6-tetramethylpiperidine)
25. 15-n-octadecyl-7-hydroxy-7,15-diazaspiro[5.1.5.3]hexadecane-14,16-dione
26. 4-benzyloxy-1-hydroxy-2,2,6,6-tetramethylpiperidine
27. 3-hydroxy-2,2,4,4-tetramethyl-20-(2-lauryloxy-carbonyl)ethyl-7-oxa-3,20-diazaspiro[5.1.11.2]heneicosan-21-one 30
28. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) phthalate
29. 1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl stearate
30. 1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl acrylate
31. 4-(4-tert-butylbenzoyloxy)-1-hydroxy-2,2,6,6-tetramethylpiperidine 35
32. (1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)4-hydroxy-3,5-di-tert-butylbenzoate
33. (1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)4-(hydroxy-3,5-di-tert-butylbenzoyl)-3,5-di-tert-butylbenzoate
34. (1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)[4-(2-oxazepin-1-yl)-2,2,6,6-tetramethylpiperidin-1-yl]acetate 40
35. alpha,alpha'-di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)-p-xylene
36. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) diethylmalonate
37. N-n-butyl-N-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)-4-hydroxy-3,5-di-tert-butylbenzamide
38. tetrakis (1-hydroxy-2,2,6,6-tetramethyl-4-piperidyl) 1,2,3,4-butanetetra-carboxylate
39. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) n-butylmalonate 45
40. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) isophthalate
41. di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) 2-(4-hydroxy-3,5-di-tert-butylbenzyl)-n-butylmalonate

The coating composition of this invention may be ambient curable or acid catalyzed hot curable systems, depending form the type of binder resins in the composition. 50

Resins for ambient curable coatings may be, for example, alkyd resins, thermoplastic acrylic resins, acrylic alkyd resins or polyester resins, or said resins modified with silicones, isocyanates, epoxides, isocyanurates, ketimines or oxazolidines. The resins may be esters of cellulose such as nitrocellulose or cellulose acetobutyrate or the resins may be epoxide resins hardenable with polyamines or other hardeners. 55

Applicable alkyd, acrylic, polyester and epoxide resins are described in S. Paul's "Surface Coatings: Science and Technology" (1985) at pages 70-310. The unmodified and modified alkyd resins which can be stabilized in accordance with the invention, are the conventional resins which are used in trade sales, maintenance and automotive refinish coatings. For example, such coatings are based on alkyd resins, alkyd/acrylic resins and alkyd/silicon resins optionally crosslinked by isocyanates or epoxy resins. 60

In addition various acrylic lacquer coating compositions are disclosed in US 4,168,249. Other acrylic/alkyd resins with polyisocyanate additives are disclosed in US 4,471,083; and acrylic resins containing either pendant amino ester groups or glycidyl groups are described in US 4,525,521. 65

Resins for acid catalyzed thermosetting coatings may be, for example, hot crosslinkable acrylic, polyester, polyurethane, polyamide or alkyd resins. This implies mixtures of those resins or mixtures with crosslinking agents such as melamine resins.

The acrylic resin lacquers are the conventional acrylic resin stoving lacquers or thermosetting resins including acrylic/melamine systems which are described, for example, in H. Kittel's "Lehrbuch der Lacke und Beschichtungen", Vol. 1. Part 2, on pages 735 and 742 (Berlin 1972), "Lackkunstharze" (1977), by H. Wagner and H.F. Sarx, on pages 229-238, and in S. Paul's "Surface Coatings: Science and Technology" (1985).

5 The polyester lacquers are the conventional stoving lacquers described e.g. in H. Wagner and H.F. Sarx, op. cit., on pages 86-99.

The alkyd resin lacquers which can be stabilized in accordance with the invention, are the conventional stoving lacquers which are used in particular for coating automobiles (automobile finishing lacquers), for example lacquers based on alkyd/melamine resins and alkyd/acrylic/melamine resins (see H. Wagner and H.F. Sarx, op. cit., pages 99-123). Other crosslinking agents include glycoluril resins, blocked isocyanates or epoxy resins.

In their industrial uses, enamels with high solids content based on crosslinkable acrylic, polyester, urethane or alkyd resins are cured with an additional acid catalyst. Light stabilizers containing a basic nitrogen group are generally less than satisfactory in this application. Formation of a salt between the acid catalyst and the light stabilizer leads to incompatibility or insolubility and precipitation of the salt and to a reduced level of cure and to reduced light protective action and poor resistance to moisture.

The N-hydroxy compounds of the present invention are less basic than the corresponding hindered amine compounds from which they are derived, yet still exhibit undiminished, indeed enhanced, light stabilization efficacy.

20 The ambient curable coatings as well as the acid catalyzed hot curable coatings stabilized in accordance with the invention are suitable both for metal finish coatings and solid shade finishes, especially in the case of retouching finishes. The lacquers stabilized in accordance with the invention are preferably applied in the conventional manner by two methods, either by the single-coat method or by the two-coat method. In the latter method, the pigment-containing base coat is applied first and a covering coat of clear lacquer applied over it.

The amount of hindered amine derivative employed is 0.1 to 10 % by weight, based on the solvent-free binder, preferably 0.5 to 5 % by weight. The binders can be dissolved or dispersed in customary organic solvents or in water or can be solvent-free.

When used in two-coat finishes, the hindered amine derivative can be incorporated in the clear coat or both 30 in the clear coat and in the pigmented base coat. In the manufacture of acrylic modified alkyd resins or acrylic resins, polymerizable hindered amine derivatives can be polymerized into the resin. The incorporation into the lacquer binder can also, however, be effected via polycondensation in the manufacture of the alkyd or polyester resins. In this cases, there is the additional advantage that the light stabilizers cannot be removed by extraction or migration so that their action is very prolonged.

35 To attain maximum light stability, the concurrent use of other conventional light stabilizers can be advantageous. Examples are UV absorbers of the benzophenone, benzotriazole, acrylic acid derivative, or oxalanilide type, or aryl-s-triazines or metal-containing light stabilizers, for example organic nickel compounds. In two-coat systems, these additional light stabilizers can be added to the clear coat and/or the pigmented base coat.

40 If such combinations are employed, the sum of all light stabilizers is 0.2 to 20 % by weight, preferably 0.5 to 5 % by weight, based on the film-forming resin.

Examples of the UV absorbers which may be used in the instant compositions in conjunction with the aforementioned hindered amine compounds are:

(a) 2-(2'-Hydroxyphenyl)-benzotriazoles, for example the 5'-methyl-, 3',5'-di-tert-butyl-, 5'-tert-butyl-, 45 5'-(1,1,3,3-tetramethylbutyl)-, 5-chloro-3',5'-di-tert-butyl-, 5-chloro-3'-tert-butyl-5'-methyl-, 3'-sec-butyl-5'-tert-butyl-, 4'-octoxy-, 3',5'-di-tert-amyl derivative.

(b) 2-Hydroxy-benzophenones, for example, the 4-hydroxy-, 4-methoxy-, 4-octoxy-, 4-decyloxy-, 4-dodecyloxy-, 4-benzyloxy, 4,2',4'-trihydroxy- and 2'-hydroxy-4,4'-dimethoxy derivative.

(c) Acrylates, for example, alpha-cyano- β , β -diphenyl-acrylic acid ethyl ester or isooctyl ester, 50 alpha-carbomethoxy-cinnamic acid methyl ester, alpha-cyano- β -methyl-p-methoxy-cinnamic acid methyl ester or butyl ester, alpha-carbomethoxy-p-methoxy-cinnamic acid methyl ester, N-(β -carbomethoxy- β -cyanovinyl)-2-methyl-indoline.

(d) Nickel compounds, for example, nickel complexes of 2,2'-thiobis-[4-(1,1,3,3-tetramethylbutyl)-phenol], such as the 1:1 or 1:2 complex, optionally with additional ligands such as n-butylamine, triethanolamine or N-cyclohexyl-di-ethanolamine, nickel dibutyldithiocarbamate, nickel salts of 4-hydroxy-3,5-di-tert-butylbenzylphosphonic acid monoalkyl esters, such as of the methyl, ethyl or butyl ester, nickel complexes of ketoximes such as of 2-hydroxy-4-methyl-phenyl undecyl ketonoxime, nickel complexes of 1-phenyl-4-lauroyl-5-hydroxy-pyrazol, optionally with additional ligands.

(e) Oxalic acid diamides, for example, 4,4'-di-octyl-oxyoxanilide, 2,2'-di-octyloxy-5,5'-di-tert-butyl-oxanilide, 2,2'-di-dodecyloxy-5,5'-di-tert-butyl-oxanilide, 2-ethoxy-2'-ethyl-oxanilide, N,N'-bis-(3-dimethylaminopropyl)-oxalamide, 2-ethoxy-5-tert-butyl-2'-ethyl-oxanilide and its mixture with 2-ethoxy-2'-ethyl-5,4'-di-tert-butyl-oxanilide, and its mixtures of ortho- and paramethoxy- as well as of o- and p-ethoxy-disubstituted oxanilides.

(f) Hydroxyphenyl-5-triazines such as 2,6-bis-(2,4-di-methylphenyl)-4(2-hydroxy-4-octyloxyphenyl)-s-triazine or the corresponding 4(2,4-dihydroxyphenyl) derivatives.

Of particular value in the instant compositions are the benzotriazoles of high molecular weight and low volatility such as 2-[2-hydroxy-3,5-di(alpha,alpha-dimethylbenzyl)-phenyl]-2H-benzotriazole, 2-(2-hydroxy-3,5-di-tert-octylphenyl)-2H-benzotriazole, 2-(2-hydroxy-3-alpha,alpha-dimethylbenzyl-5-tert-octylphenyl)-2H-benzotriazole, 2-(2-hydroxy-3-tert-octyl-5,alpha,alpha-dimethylbenzylphenyl)-2H-benzotriazole, 2-(2-hydroxy-3,5-di-tert-amylphenyl)-2H-benzotriazole, 2-[2-hydroxy-3-tert-butyl-5-(2-omega-hydroxy-octa(ethyleneoxy)-carbonyl)-ethylphenyl]-2H-benzotriazole, 2-[2-hydroxy-3-tert-butyl-5-(2-octyloxycarbonyl-ethyl)-phenyl]-benzotriazole, 2-(2-hydroxy-3-sec.dodecyl-5-methyl-phenyl)-benzotriazole, hexamethylene di[β-(3-tert-butyl-4-hydroxy-5-[2-benzotriazolyl]-phenyl)-propionate] and the 5-chloro compounds corresponding to each of the above named benzotriazoles.

A preferred embodiment of the instant invention pertains to stabilized compositions comprising

- (a) a resin system as defined hereinabove,
- (b) an N-hydroxyl substituted 2,2,6,6-tetraalkylpiperidine compound,
- (c) an UV absorber selected from the group consisting of the benzophenones, benzotriazoles, acrylic acid derivatives, organic nickel compounds, aryl-s-triazines and oxanilides.

Further ingredients which the enamels or coatings can contain are antioxidants, for example those of the sterically hindered phenol derivative type, phosphorus compounds, such as phosphites, phosphines or phosphonites, conventional hindered amine light stabilizers, plasticizers, levelling assistants, hardening catalysts, thickeners, dispersants or adhesion promoters.

A further preferred embodiment of the instant invention is a stabilized composition containing components (a), (b) and (c) described above which additionally contains as component (d) a phosphite or phosphonite.

The amount of phosphite or phosphonite (d) which is used in the instant compositions is from 0.05 to 2 % by weight, preferably from 0.1 to 1 % by weight, based on the film forming resin. In two-coat systems, these stabilizers may be added to the clear coat and/or base coat.

Typical phosphite and phosphonites include triphenyl phosphite, diphenylalkyl phosphites, phenyldialkyl phosphites, tri-(nonylphenyl)-phosphite, triauryl phosphite, trioctadecyl phosphite, di-stearyl-pentaerythritol diphosphite, tris-(2,4-di-tert-butylphenyl) phosphite, di-isodecyl-pentaerythritol diphosphite, di-(2,4-di-tert-butylphenyl)-pentaerythritol diphosphite, tristearyl-sorbitol triphosphite, tetrakis-(2,4-di-tert-butylphenyl)-4,4'-di-phenylenediphosphonite.

The stabilizers are needed to impart greater retention of durability to the cured enamels (as measured by 20° glass, distinction of image, cracking or chalking); the enamel should not yellow on curing and further color change on exposure to light should be minimized; the stabilizers should be soluble in the organic solvents normally used in coating applications such as methyl amyl ketone, xylene, n-hexyl acetate, alcohol and the like.

The instant hindered amine light stabilizers substituted on the hindered N-atom by a hydroxy moiety fulfill each of these requirements and provide individually or in combination with a UV-absorber outstanding light stabilization protection to the cured coatings.

The following examples describe the inventive use of substituted hindered amine derivatives in various ambient curable and acid catalyzed hot curable resin systems. Parts and percentages are by weight.

Example 1: Stabilization of a White, Two-Component Acrylic Urethane Gloss Enamel

A white acrylic urethane enamel is formulated as shown below.

Component I	Parts
Acryloid® AU 608	83.5
(acrylic polyol from Rohm & Haas)	
TiO ₂	196.7
Cellosolve Acetate	97.3
- Sand Mill -	
Acryloid® AU 608	291.4
Flow Improver	0.28
Cellosolve Acetate	155.5
Component II	
Desmodur® N-100	70.9
(polyisocyanate from Mobay Corp)	
Cellosolve Acetate	101.7
	997.3

This material is spray applied at a dry film thickness of 35-50 μm onto Bonderite 40 cold rolled steel panels that had been previously primed with a commercial epoxy polyamide maintenance primer (Sherwin-Williams Tile Clad II). Prior to application, the indicated amount of additive (based on resin solids) is added to the paint.

After ambient storage for a period of 2 weeks, the coated panels are subjected to weathering for 1790 hours

in a QUV tester. In this test, the samples are subjected to weathering for 8 hours in a humid atmosphere and UV light at 50°C and for 4 hours with no UV light at 40°C. The 60° gloss of each panel (ASTM D-523) is determined and is reported below.

5	<u>Additive</u>	<u>Conc. (% by wt.)</u>	<u>60° Gloss</u>
	A	1.0	27
	Compound	1.0	48
10	No. 5		
	A = bis(1,2,2,6,6-pentamethyl-4-piperidinyl)sebacate		

15

Example 2: Stabilization of Semi-Transparent Alkyd Stain

A semi-transparent linseed alkyd stain is formulated as shown below.

20	<u>Materials</u>	<u>Parts</u>
	Copolymer 186 (dehydrated castor oil from CasChem)	27.3
25	Mineral Spirits	36.0
	Lecithin	4.0
	Bentone SD-1	10.0
	Yellow Iron Oxide	4.36
	Red Iron Oxide	19.58
30	Black Iron Oxide	3.3
	Fungitrol® 11	2.0
	- Cowles Grind -	
	Copolymer 186	66.3
35	Beckosol® 10-051 (long oil alkyd from Reichold)	115.0
	Linseed Oil	20.3
	Mineral Spirits	118.7
	6 % Cobalt	1.2
40	Naphthenate	
	5 % Calcium	2.16
	Naphthenate	
	18 % Zirconium	1.2
	Octoate	
45	Ketoxime (antiskinning agent)	1.0
	Xylol	21.6
	Mineral Spirits	273.4
50		<hr/> 726.7

Pieces of 1.3 cm. x 20.3 cm x 30.5 cm Western Red Cedar panels having a fine radial cut are used. One half of each panel is coated with 3 coats of the unstabilized stain. An equal amount of stain containing 3 % (by weight on resin solids) of light stabilizers is applied to the other half of the panel in three coats. After storage at ambient temperatures for two weeks, the wood panels are exposed outdoors vertically facing south for a period of eight months. The 60° gloss retention of the stabilized and unstabilized section of the same panel are determined. Due to lack of homogeneity of wood substrates, the gloss retention of the same stain tends to differ from panel to panel. In this study, the unstabilized control stain is applied to every panel inasmuch as the differences in gloss retention between the stabilized and unstabilized portions of the panel allow for a better measure of the improvement in gloss due to the presence of the light stabilizer.

65

Additive	60° Gloss Value		
	Unsta- bilized	Stabilized	Gloss Improve- ment
3 % A	21.4	20.3	-
3 % 29	27.5	69.8	42.3
1.5 % B + -	16.9	32.4	15.5
1.5 % A			
1.5 % B + -	23.9	66.7	42.8
1.5 % 29			

B = 5-(alpha,alpha-dimethylbenzyl)-2-(2-hydroxy-3,5-di-tert.butylphenyl)-2H-benzotriazole

Example 3: Stabilization of a Medium Oil Alkyd Enamel

A commercially available medium oil alkyd enamel pigmented with non-leaving aluminium pigment and tinted light blue is stabilized with the indicated amount of UV absorber and hindered amine derivative (by weight of resin solids) and then spray applied onto cold rolled steel panels primed with an epoxy primer. After the coating is allowed to cure at room temperature for 2 weeks, the panels are exposed in a Xenon Arc Weatherometer for 840 hours. The 20° gloss retention (ASTM D-523-80) of the panels is determined.

Additive	Conc. (% by wt.)	20° Gloss Retention %
C/A	3/2	22.7
C/29	3/2	34.4
C/39	3/2	31.9

C = 2-[2-hydroxy-3-tert.butyl-5-(2-omega-hydroxy-octa-(ethyleneoxy)carbonylethylphenyl)]-2H-benzotriazole

Example 4: Stabilization of a Nitrocellulose Lacquer

A commercially available white nitrocellulose lacquer (from Randolph Corp.) is stabilized with the indicated light stabilizers (by weight of resin solids) and spray applied onto epoxy primed cold rolled steel panels. After storage at room temperature for 2 weeks, the panels are exposed in a Xenon Arc Weatherometer for 1130 hours. The 60° gloss of the panels at that interval is determined.

Additive	Conc. (% by wt.)	60° Gloss
C/A	1.5/1.5	41
C/39	1.5/1.5	70
D/A	1.5/1.5	36
D/39	1.5/1.5	70

C = 2-(2'-hydroxy-3'-(1-methylundecyl)-5'-methyl)-2H-benzotriazole

Example 5: Stabilization of a Blue Medium Oil Alkyd Enamel

A blue medium oil alkyd enamel is formulated as follows:

EP 0 309 401 A1

	<u>Material</u>	<u>Wt. %</u>
	Aroplaz® 1445-M-50 (alkyd resin solution from NL Industries)	66.6
5	TiO ₂	2.5
	Phthalocyanine Blue	0.8
	Driers (Co, Pb, Mn)	1.0
	Xylene	<u>29.1</u>
10		100.0

15 This material is spray applied to a thickness of 45 µm onto Bonderite 40 cold rolled steel panels which have been previously electrocoated with an epoxy ester (PPG Uniprine). The panels are allowed to cure at room temperature for 2 weeks and are then exposed outdoors at an angle of 45° facing south. After 9 months exposure, the 20° gloss retention of the coating is determined.

	<u>Additive</u>	<u>Conc. (% by wt.)</u>	<u>20° Gloss Retention %</u>
20	Control (unstabilized)	-	0
	E/5	2/1	69
25	E = 2-(2'-hydroxy-3',5'-di-tert.butylphe- nyl)benztriazole		

30 Example 6: The following alkyd paint formulations are prepared.

35

40

45

50

55

60

65

	White(W)	Parts Yellow(Y)	Blue(B)	
Aroplaz® 1445 M-50	45.0	57.73	58.27	5
TiO ₂	45.0	20.82	4.26	
Irgalite®	-	7.3	-	
GS (yellow pigment from CIBA- GEIGY)				10
Phthalocy- anine blue	-	-	1.41	
Ketoxime (antiskin- ning agent)	0.17	0.07	0.014	15
Ionic antifloat compound	-	-	0.05	20
24 % Pb as naphthe- nate	0.94	1.30	1.30	
6 % Co as naphthe- nate	0.36	0.50	0.50	25
6 % Mn as naphthe- nate	0.45	0.60	0.60	30
Xylene	35.0	48.4	45.1	
Mineral Spirits	60.0	64.49	64.11	35

The formulations are stabilized with the indicated materials in the indicated concentrations (by weight on total resin solids) and sprayed onto cold rolled steel panels primed with an electrocoated epoxy primer. The coating is allowed to cure overnight at room temperature and the panels are then exposed in Florida at an angle of 45° South. 60° gloss, distincton of image (DI) (Hunter Associates Apparatus) and color change based on Yellowness Index values are determined and tabulated below.

Additive	Conc. (% by wt.)	Paint	Florida Exposure (Months)									
			60° Gloss			DI			Color Change			
			0	9	12	15	0	9	12	15	12	15
-	-	W	82	35`	33	35	96	15	35	18	4.7	6.3
E/5	2/1	W	82	55	49	45	96	74	66	53	3.5	5.2
-	-	Y	92	16	13	10	95	5	6	2	14	14
E/5	2/1	Y	92	60	56	33	95	85	76	43	10	12
-	-	B	91	26	17*	-	93	15	25	-	10	-
E/5	2/1	B	91	84	75	54	93	82	78	68	2	3.4

* = cracking observed

Example 7: Stabilization of an Acrylic Alkyd Enamel Crosslinked with an Aliphatic Isocyanate Refinish Enamel

A commercially available light blue metallic acrylic alkyd enamel hardened with an aliphatic isocyanate is stabilized with the indicated amounts of additive (by weight on total resin solids) and then spray applied onto Bonderite 40 panels primed with an alkyd primer. After the coatings are aged at ambient temperature for 2 weeks, the panels are exposed outdoors at an angle of 5° for a period of 10 months. The 20° gloss retention of the panels is measured.

	Additive	Conc. (% by wt.)	20° Gloss Retention %
	C/F	3/2	16
5	C/36	3/2	27
	C/40	3/2	25
	C/39	3/2	24
	C/28	3/2	22

10 F = bis(1,2,2,6,6-pentamethyl-4-piperid-
inyl)[[3,5-di-tert.-butylhydroxyphenyl]methyl]bu-
tylpropanedioate

15

Example 8: Stabilization of a Thermoplastic Acrylic Lacquer

A commercially available light blue metallic thermoplastic acrylic lacquer is stabilized with the indicated amounts of additive (by weight on total resin solids) and then spray applied onto Bonderite 40 panels primed with an epoxy primer. After storage at ambient temperature for 2 weeks, the panels are exposed in a Xenon
20 Arc Weatherometer for 1250 hours. The 60° gloss of the panels is determined.

	Additive	Conc. (% by wt.)	60° Gloss
25	G/A	2/2	11
	G/5	2/2	27
	G/32	2/2	24
	G/29	2/2	27
	G/28	2/2	31
30	G/41	2/2	23
	G/39	2/2	39
	G/40	2/2	23
	G/36	2/2	23
35	G/38	2/2	26

G = 2-[2-hydroxy-3,5-di
(alpha,alpha-dimethylbenzyl)-phenyl]-2H-benzo-
triazole

40

Example 9: Stabilization of an Acrylic Alkyd Refinish Enamel

45 A commercially available acrylic enamel pigmented with non-leafing aluminium pigment and tinted a light blue is stabilized with the indicated amounts of UV absorber and hindered amine (by weight of resin solids) and then spray applied onto Bonderite 40 panels primed with an alkyd primer. After the coating is allowed to cure at room temperature for 14 days, the panels are exposed outdoors at an angle of 5° S for a period of 10 months. The 20° gloss retention of the panels is measured.

	Additive	Conc. (% by wt.)	20° Gloss
50	C/A	3/2	18
	C/5	3/2	25
55	C/28	3/2	26
	C/39	3/2	25
	C/40	3/2	28

60

Example 10: Stabilization of High Solids Acid-catalyzed, Thermoset Acrylic Resin Enamel

A high solids (50 % by weight) thermoset acrylic resin enamel, catalyzed by 0.5 % by weight of p-toluenesulfonic acid, based on the film-forming resin, is stabilized by the addition of various N-hydroxy-polyalkylpiperidine derivatives.

65 The high solids thermoset acrylic resin enamel formulation (Acryloid® AT 400 from Rohm & Haas) is based

on hydroxyethyl methacrylate, methyl methacrylate, styrene, butyl acrylate and butyl methacrylate and a melamine curing agent.

Pieces of steel sheeting 9 x 30 cm, coated with a primer based on polyester/epoxy resin, are then coated with a silver metallic base coat and finally with the clear finishing enamel. The base coat is sprayed onto the sheet to a thickness of about 20 µm and air dried for 3 minutes. The clear finishing enamel coat is then sprayed onto the sheet to a thickness of about 45 µm. After 15 minutes air-drying, the coated sheets are baked for 30 minutes at 250°F (121°C).

The stabilizers under test are added to the thermoset acrylic resin finishing enamel before the enamel is sprayed onto the base coated sheet.

After storage for 3 weeks in an air-conditioned room (23°C/50 % relative humidity), the coated sheets are subjected to weathering for 1300 hours according to the method ASTM G-53/77 in a QUV tester. In this test, the samples are subjected to weathering in repeated cycles for 4 hours in a humid atmosphere at 50°C and then for 8 hours under UV light at 70°C. Subsequently, distinction of the image (DI) and the 20° gloss of the surface of the finishes are determined. High values for 20° gloss and DI are desired.

QUV Exposure - 1300 hours

<u>Stabilizer in clear coat</u>	<u>20° Gloss</u>	<u>DI</u>
Without Additive	6	12
1 % Compound G + 2 % Compound 5	92	66
1 % Compound G + 2 % Compound 12	90	68

Example 11:

In another test, the samples, prepared as in Example 10, after storage are subjected to accelerated weathering for 2230 hours in a Xenon arc (6500 Watt) Weather-Ometer.

The samples are exposed to repeated cycles for 102 minutes to the light source, then for 18 minutes to a water spray and light, at 60°C (black panel). The results are reported below. High values for 20° gloss and DI are desired.

Xenon arc exposure

<u>Stabilizer in clear coat</u>	<u>20° Gloss</u>	<u>DI</u>
Without Additive	44	6
1 % Compound G + 2 % Compound 5	89	70
1 % Compound G + 2 % Compound 12	89	84

Example 12: Cracking Resistance

Samples described in Example 10 after being exposed in the QUV tester for 1300 hours are evaluated visually for cracking. The results are given below.

Cracking after 1300 hours

	<u>Stabilizer in clear coat</u>	<u>Appearance of the Enamel</u>
5	Without Stabilizer	Deep cracks
	1 % Compound G + 2 % Compound 5	No cracking
	1 % Compound G + 2 % Compound 12	No cracking
10		

Example 13: Knoop Hardness and Enamel Appearance

15 Samples described in Example 10 after baking for 30 minutes at 121°C are stored for one week at room temperature. Knoop hardness values are determined on the baked enamel and the general appearance of the enamels is noted.

	<u>Stabilizer in clear coat</u>	<u>Baked Coating Knoop Hardness</u>	<u>Appearance of Coating Before Baking</u>
20	Without Stabilizer	11.8	Clear
25	1 % Compound G + 2 % Compound 5	11.6	Clear
30	1 % Compound G + 2 % Compound 12	10.6	Clear

35

Example 14:

Two thermoset acrylic enamels are formulated to include a benzotriazole UV-absorber and a hindered amine light stabilizer as described in Example 10.

40 The coated panels are exposed in the QUV tester, or in the Xenon Arc Weatherometer or in South Florida in an unheated black box at an angle of 5° to the sun. 20° gloss or distinction of image (DI) values are determined.

The thermoset acrylic enamels are based on a binder of 70 % of monomers such as hydroxyethyl acrylate, styrene, acrylonitrile, butyl acrylate and acrylic acid with 30 % of a melamine resin and as acid catalyst 0.5 % of p-toluene-sulfonic acid.

45 Enamel A is coated over a silver metallic base coat and Enamel B is coated over a white base coat.

50

55

60

65

20° Gloss Values after Hours of QUV Exposure

<u>Enamel A with</u>	<u>0</u>	<u>750</u>	<u>1300</u>	<u>1600</u>	<u>2350</u>	<u>3000</u>	<u>Hours of QUV Exposure Till First Cracking Observed</u>	
No Stabilizer	95	82	8	2	-	-	900	5
1 %	92	92	90	90	40	10	2700	10
Compound G plus 2 %								15
Compound 5								20

20° Gloss Values after Hours of Xenon Arc Exposure

<u>Enamel A with</u>	<u>0</u>	<u>1400</u>	<u>1850</u>	<u>2500</u>	<u>2750</u>	<u>3300</u>	<u>4050</u>	<u>5100</u>	<u>5900</u>	<u>7400</u>	<u>Hours of Xenon Arc Exposure Till First Cracking Observed</u>	
No Stabilizer	90	55	35	22	15	0	-	-	-	-	2700	25
1 %	90	90	90	90	90	88	88	75	70	45	> 8500	30
Compound G plus 2 %												35
Compound 5												40

Distinction of Image Values after Months of South Florida Exposure

<u>Enamel A with</u>	<u>0</u>	<u>6</u>	<u>12</u>	<u>18</u>	
No Stabilizer	82	25	5	2	45
1 %	85	82	75	60	
Compound G plus 2 %					50
Compound 5					

Distinction of Image Values after Hours of QUV Exposure

<u>Enamel B with</u>	<u>0</u>	<u>200</u>	<u>1050</u>	<u>1450</u>	<u>1700</u>	<u>1840</u>	<u>2250</u>	
No Stabilizer	78	55	8	5	-	-	-	55
1 %	88	88	88	88	88	88	60	
Compound G plus 2 %								60
Compound 5								65

Example 15:

In order to determine whether cure is affected at either the 121°C normal bake temperature or at the 82°C automotive low bake repair temperature, Enamel A (as described in Example 14) containing various stabilizer combinations is baked over a silver base coat for 30 minutes at 121°C or for 30 minutes at 82°C.

- 5 The effectiveness of cure is assessed from Knoop hardness values, the higher numbers indicating greater hardness and hence cure.

	<u>Enamel A containing 1 % by weight Compound G plus 2 % by weight of Test Compound</u>	<u>Knoop Hardness Values</u>	
		<u>Normal Bake</u>	<u>Low Bake Repair</u>
10			
	Unstabilized	11.5	11.5
	Compound A*)	1-2	-
20	Compound 5	11.5	7.0

* Compound A is
di-(1,2,2,6,6-pentamethylpiperidin-4-yl) sebacate.

- 25 The instant compound 5 does not retard cure at the normal bake temperature and causes only slight insignificant retardation of cure at the automotive low bake repair temperature.

Example 16:

- 30 The physical properties of Enamel A (as described in Example 14) coated over a silver metallic base coat when cured for 30 minutes at 82°C, the low bake repair temperature, are completely satisfactory as is seen in the table below.

	<u>Stabilizer</u>	<u>Knoop Hardness</u>	<u>Pendulum Hardness</u>	<u>Change in 20° Gloss Value after Humidity</u>	<u>Crosshatch Adhesion</u>
35					
	Unstabilized	7	70	20	4
	1.5 % 5 + 1 % G	7	85	2	5

- 40 The 20° gloss values are taken on coated panels conditioned for 96 hours in a constant humidity chamber at 38°C and 100 % relative humidity. A low change in 20° gloss value shows the coating is not adversely affected by humidity.

- 45 The cross-hatch type adhesion test involves using a multi-cut knife to prepare cross-hatches through the topcoat film and base coat on the panel. An acetate fiber adhesive tape is placed over the cross-hatch area and then is pulled off. A visual inspection of the amount of topcoat, if any, coming off with the tape as it is pulled gives a relative rating of the amount of delamination. A rating system of 5 indicating the absence of cross-hatch delamination to 0 indicating complete cross-hatch delamination is used.

- 50 It is clear that the low bake curing at 82°C for the acid catalyzed thermoset enamels containing the N-hydroxy hindered amine light stabilizers is completely adequate.

Example 17:

- 55 The thermoset acrylic enamel of Example 14 including 0.5 % of p-toluenesulfonic acid is formulated to include varying concentrations of a benzotriazole UV-absorber and a hindered amine light stabilizer of the invention. The enamel is coated over a silver metallic base coat pursuant to the procedure in Example 14 and baking is conducted for 30 minutes at 121°C normal bake temperature.

The coated panels are exposed in the QUV exposure apparatus and the time to 50 % loss of 20° gloss is determined.

60

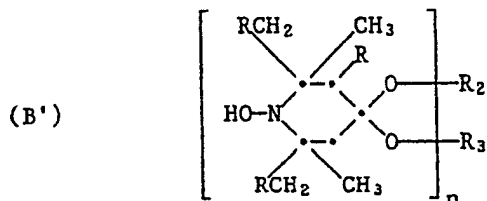
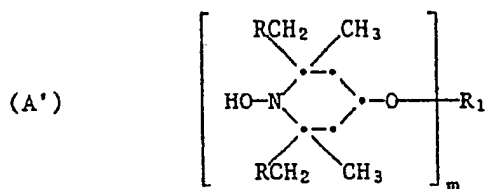
65

Compound	Conc. (% by wt.)	Time to 50 % Loss of 20° Gloss (hours)
unstabilized	-	600
G/5	2/1.5	3300
G/22	2/1.5	2900
G/29	2/1.5	2000
G/37	2/1.5	1700
unstabilized	-	900
C/5	3.5/1.5	4750
C/5	3.5/1.5	5300
C/28	3.5/1.5	4500
C/29	3.5/1.5	5300
C/31	3.5/1.5	3900
C/35	3.5/1.5	4700
C/39	3.5/1.5	5100
C/40	3.5/1.5	5300
unstabilized	-	1000
C/5	3/1	4700
C/27	3/1	4100
C/36	3/1	5200
C/38	3/1	4600
C/39	3/1	4200
C/40	3/1	4600

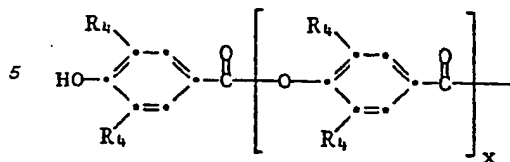
Various N-hydroxy derivatives of hindered amines are known. For example, a substantial number of such derivatives are generically disclosed in US 4,590,231. The substituents on the 4-position include ether, ester, amino, dioxaspiro, diaza-dione, urea and triazine substituents, among others. These derivatives are identified as stabilizers in polyolefin compositions. Mono- and di-piperidinyl ester derivatives are likewise disclosed in JP-A 54-69162 for use in stabilizing urethane polymers. Other N-hydroxy derivatives are disclosed in J. Polym. Sci., Polymer Chem. Ed. 22, 277-281 (1984); Polym. Sci. Technol. 26, 35-47 (1984); Zh. Org. Khim. 6, 2365-2369 (1970); as well as in US 3,936,456, US 4,404,302 and US 4,472,547.

Accordingly, it is a further object of the instant invention to identify a series of new N-hydroxy hindered amines having a broad range of stabilization performance characteristics.

These derivatives thus correspond to the formulae A' or B'

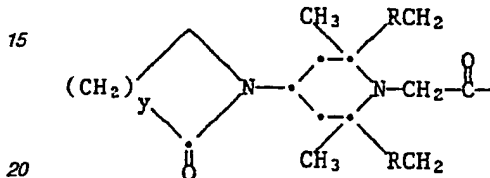


wherein R is hydrogen or methyl, preferably is hydrogen;
m is 1-4;
when m is 1, R₁ is



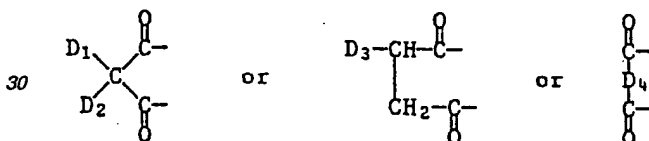
10

wherein x is 0 or 1 and R₄ independently are C₁-C₈alkyl, or



with y is 2-4;
when m is 2, R₁ is

25



35 wherein D₁ and D₂ are independently C₁-C₈alkyl, C₆-C₁₀aryl, C₇-C₁₂aralkyl or 3,5-di-*t*-butyl-4-hydroxybenzyl and D₂ is also hydrogen, D₃ is C₁-C₁₈alkyl or C₂-C₁₈alkenyl, and D₄ is C₅-C₁₂cycloalkylene;

when m is 3, R₁ is a trivalent acyl radical of an aliphatic, unsaturated aliphatic, cycloaliphatic or aromatic tricarboxylic acid having up to 20 C-atoms;

when m is 4, R₁ is a tetravalent acyl radical of a saturated or unsaturated aliphatic or aromatic tetracarboxylic acid having up to 20 C-atoms;

40

n is 1 or 2;

when n is 1, R₂ and R₃ are independently hydrogen, C₁-C₁₂alkyl, C₂-C₁₂alkenyl or C₇-C₁₂aralkyl; and

when n is 2, R₂ and R₃ together are (-CH₂)₂C(CH₂)₂.

45 In these formulae, representative alkyl groups include methyl, ethyl, *n*-propyl, *n*-butyl, *tert*-butyl, *n*-hexyl, *n*-octyl, 2-ethylhexyl, *n*-undecyl and *n*-dodecyl; aryl includes phenyl and naphthyl; aralkyl includes benzyl, alpha-methylbenzyl and phenethyl; and alkenyl includes 1-propenyl, allyl, methallyl, 2-butenyl, 2-hexenyl and 2-octenyl. R₁ as a trivalent acyl radical is, for example, an acyl radical of benzene-1,2,4-tricarboxylic acid. R₁ as a tetravalent acyl radical is, for example, an acyl radical of 1,2,3,4-butanetetracarboxylic acid, 1,2,3,4-but-2-enetetracarboxylic acid and 1,2,3,5- and 1,2,4,5-pentanetetracarboxylic acid. In the R₁ members when m is 2, malonates are preferred where D₁ is C₁-C₈alkyl or 3,5-di-*t*-butyl-4-hydroxybenzyl and D₂ is D₁ or hydrogen.

50

Preferred are the compounds of formula A' wherein

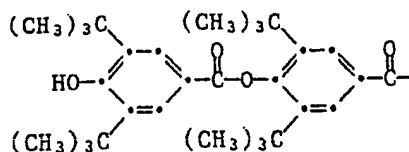
R is hydrogen, m is 1, 2 or 4 and

when m is 1, R₁ is a group

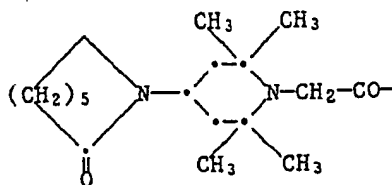
55

60

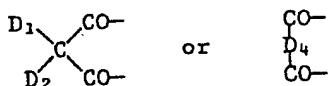
65



or



when m is 2, R_1 is a group



wherein D_1 is C_1 - C_4 alkyl or 3,5-di- t -butyl-4-hydroxybenzyl and D_2 is D_1 or hydrogen and D_4 is cyclohexylene, and when m is 4, R_1 is the tetravalent acyl radical of a butanetetracarboxylic acid.

The N-hydroxy derivatives of formula A' and B' are generally prepared by oxidizing the corresponding N-H hindered amine with an appropriate peroxy compound such as hydrogen peroxide or tert-butyl hydroperoxide in the presence of molybdenum oxide or a metal carbonyl or metal oxide catalyst followed by reduction of the oxyl intermediate formed thereby to the desired N-hydroxy derivative, preferably by catalytic hydrogenation. Molybdenum (VI) is noted to increase the efficiency of the oxidation step. The reaction is preferably conducted in hydrocarbon solvents such as toluene. The starting materials needed to prepare the derivatives of this invention are items of commerce or can be prepared by known methods.

The derivatives are particularly effective in stabilizing organic materials primarily against the degradative effects of actinic stimuli. Such organic materials include polymeric materials such as the following polymers.

1. Polymers of monoolefins and diolefins, for example polypropylene, polyisobutylene, polybutene-1, polymethylpentene-1, polyisoprene or polybutadiene, as well as polymers of cycloolefins, for instance of cyclopentene or norbornene, polyethylene (which optionally can be crosslinked), for example high density polyethylene (HDPE), low density polyethylene (LDPE) and linear low density polyethylene (LLDPE).

2. Mixtures of the polymers mentioned under 1), for example mixtures of polypropylene with polyisobutylene, polypropylene with polyethylene (for example PP/HDPE, PP/LDPE) and mixtures of different types of polyethylene (for example LDPE/HDPE).

3. Copolymers of monoolefins and diolefins with each other or with other vinyl monomers, such as, for example, ethylene/propylene, linear low density polyethylene (LLDPE) and its mixtures with low density polyethylene (LDPE); propylene/butene-1, ethylene/hexene, ethylene/ethylpentene, ethylene/heptene, ethylene/octene, propylene/isobutylene; ethylene/butene-1, propylene/butadiene, isobutylene/isoprene, ethylene/alkyl acrylates, ethylene/alkyl methacrylates, ethylene/vinyl acetate or ethylene/acrylic acid copolymers and their salts (ionomers) and terpolymers of ethylene with propylene and a diene, such as hexadiene, dicyclopentadiene or ethylidene-norbornene; as well as mixtures of such copolymers and their mixtures with polymers mentioned in 1) above, for example polypropylene/ethylene-propylene-copolymers, LDPE/EVA, LDPE/EAA, LLDPE/EVA and LLDPE/EAA.

3a. Hydrocarbon resins (for example C_5 - C_9) and hydrogenated modifications thereof (for example tackifiers).

4. Polystyrene, poly-(p -methylstyrene), poly-(α -methylstyrene).

5. Copolymers of styrene or α -methylstyrene with dienes or acrylic derivatives, such as, for example, styrene/butadiene, styrene/acrylonitrile, styrene/alkyl methacrylate, styrene/maleic anhydride, styrene/butadiene/ethyl acrylate, styrene/acrylonitrile/methyl acrylate; mixtures of high impact strength from styrene copolymers and another polymer, such as, for example, from a polyacrylate, a diene polymer or an ethylene/propylene/diene terpolymer; and block copolymers of styrene, such as, for example, styrene/butadiene/styrene, styrene/isoprene/styrene, styrene/ethylene/butylene/styrene or styrene/ethylene/propylene/styrene.

6. Graft copolymers of styrene or α -methylstyrene such as, for example, styrene on polybutadiene, styrene on polybutadiene-styrene or polybutadiene-acrylonitrile; styrene and acrylonitrile (or methacry-

lonitrile) on polybutadiene; styrene and maleic anhydride or maleimide on polybutadiene; styrene, acrylonitrile and maleic anhydride or maleimide on polybutadiene; styrene, acrylonitrile and methyl methacrylate on polybutadiene, styrene and alkyl acrylates or methacrylates on polybutadiene, styrene and acrylonitrile on ethylene/propylene/diene terpolymers, styrene and acrylonitrile on polyacrylates or polymethacrylates, styrene and acrylonitrile on acrylate/butadiene copolymers, as well as mixtures thereof with the copolymers listed under 5), for instance the copolymer mixtures known as ABS-, MBS-, ASA- or AES-polymers.

7. Halogen-containing polymers, such as polychloroprene, chlorinated rubbers, chlorinated or sulfochlorinated polyethylene, epichlorohydrin homo- and copolymers, polymers from halogen-containing vinyl compounds, as for example, polyvinylchloride, polyvinylidene chloride, polyvinyl fluoride, polyvinylidene fluoride, as well as copolymers thereof, as for example, vinyl chloride/vinylidene chloride, vinyl chloride/vinyl acetate or vinylidene chloride/vinyl acetate copolymers.

8. Polymers which are derived from α,β -unsaturated acids and derivatives thereof, such as polyacrylates and polymethacrylates, polyacrylamide and polyacrylonitrile.

9. Copolymers from the monomers mentioned under 8) with each other or with other unsaturated monomers, such as, for instance, acrylonitrile/butadiene, acrylonitrile/alkyl acrylate, acrylonitrile/alkoxyalkyl/acrylate or acrylonitrile/vinyl halogenide copolymers or acrylonitrile/alkyl methacrylate/butadiene terpolymers.

10. Polymers which are derived from unsaturated alcohols and amines, or acyl derivatives thereof or acetals thereof, such as polyvinyl alcohol, polyvinyl acetate, polyvinyl stearate, polyvinyl benzoate, polyvinyl maleate, polyvinyl butyral, polyallyl phthalate or polyallyl melamine; as well as their copolymers with olefins mentioned in 1) above.

11. Homopolymers and copolymers of cyclic ethers, such as polyalkylene glycols, polyethylene oxide, polypropylene oxide or copolymers thereof with bis-glycidyl ethers.

12. Polyacetals, such as polyoxymethylene and those polyoxymethylenes which contain ethylene oxide as a comonomer; polyacetals modified with thermoplastic polyurethanes, acrylates or MBS.

13. Polyphenylene oxides and sulfides, and mixtures of polyphenylene oxides with polystyrene or polyamides.

14. Polyurethanes which are derived from polyethers, polyesters or polybutadienes with terminal hydroxyl groups on the one side and aliphatic or aromatic polyisocyanates on the other side, as well as precursors thereof (polyisocyanates, polyols or prepolymers).

15. Polyamides and copolyamides which are derived from diamines and dicarboxylic acids and/or from aminocarboxylic acids or the corresponding lactams, such as polyamide 4, polyamide 6, polyamide 6/6, 6/10, 6/9, 6/12 and 4/6, polyamide 11, polyamide 12, aromatic polyamides obtained by condensation of m-xylene diamine and adipic acid; polyamides prepared from hexamethylenediamine and isophthalic or/and terephthalic acid and optionally an elastomer as modifier, for example poly-2,4,4'-trimethylhexamethylene terephthalamide or poly-m-phenylene isophthalamide. Further copolymers of the aforementioned polyamides with polyolefins, olefin copolymers, ionomers or chemically bonded or grafted elastomers; or with polyethers, such as for instance, with polyethylene glycols, polypropylene glycols or polytetramethylene glycols. Polyamides or copolyamides modified with EPDM or ABS. Polyamides condensed during processing (RIM-polyamide systems).

16. Polyureas, polyimides and polyamide-imides.

17. Polyesters which are derived from dicarboxylic acids and diols and/or from hydroxycarboxylic acids or the corresponding lactones, such as polyethylene terephthalate, polybutylene terephthalate, poly-1,4-dimethylolcyclohexane terephthalate, poly-[2,2-(4-hydroxyphenyl)-propane] terephthalate and polyhydroxybenzoates as well as block-copolyether-esters derived from polyethers having hydroxyl end groups.

18. Polycarbonates and polyester-carbonates.

19. Polysulfones, polyether-sulfones and polyether-ketones.

20. Crosslinked polymers which are derived from aldehydes on the one hand and phenols, ureas and melamines on the other hand, such as phenol/formaldehyde resins, urea/formaldehyde resins and melamine/formaldehyde resins.

21. Drying and non-drying alkyd resins.

22. Unsaturated polyester resins which are derived from copolyesters of saturated and unsaturated dicarboxylic acids with polyhydric alcohols and vinyl compounds as crosslinking agents, and also halogen-containing modifications thereof of low inflammability.

23. Thermosetting acrylic resins, derived from substituted acrylic esters, such as epoxy-acrylates, urethane-acrylates or polyester-acrylates.

24. Alkyd resins, polyester resins or acrylate resins in admixture with melamine resins, urea resins, polyisocyanates or epoxide resins as crosslinking agents.

25. Crosslinked epoxide resins which are derived from polyepoxides, for example from bis-glycidyl ethers or from cycloaliphatic diepoxides.

26. Natural polymers, such as cellulose, rubber, gelatine and derivatives thereof which are chemically modified in a polymer-homologous manner, such as cellulose acetates, cellulose propionates and cellulose butyrates, or the cellulose ethers, such as methylcellulose; rosins and their derivatives.

27. Mixtures of polymers as mentioned above, for example PP/EPDM, Polyamide 6/EPDM or ABS, PVC/EVA, PVC/ABS, PVC/MBS, PC/ABS, PBTP/ABS, PC/ASA, PC/PBT, PVC/CPE, PVC/acrylates, POM/thermoplastic PUR, PC/thermoplastic PUR, POM/acrylate, POM/MBS, PPE/HIPS, PPE/PA 6.6 and copolymers, PA/HDPE, PA/PP, PA/PPE.

The compounds of formula A' and B' alone or in combination with phenols can further be used in photographic layers as yellow dye light stabilizers, as cyan dye dark stabilizers, as antistain agents in magenta layers (especially for two-equivalent magenta couplers) and as thermal stabilizers for magenta couplers.

The following examples illustrate the preparation of these compounds.

Example 18: Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) methylmalonate

A 4.5M solution of t-butyl hydroperoxide in toluene (25 ml, 113 mmol) is rapidly added to a mixture of 50.0 g (126 mmol) of di-(2,2,6,6-tetramethylpiperidin-4-yl) methylmalonate, 3.0 g of molybdenum trioxide, and 100 ml of toluene at 90°C. Another portion of 4.5M t-butyl hydroperoxide in toluene (87 ml, 392 mmol) is added over 20 minutes. The reaction mixture is stirred for 3 hours at 85-90°C, then cooled and filtered. The filtrate is concentrated and crystallized from methanol to yield 20.5 g (38 %) of di-(1-oxyl-2,2,6,6-tetramethylpiperidin-4-yl) methylmalonate, an orange solid, m.p. 111-120°C.

A mixture of 20.3 g (47.6 mmol) of di-(1-oxyl-2,2,6,6-tetramethylpiperidin-4-yl) methylmalonate, 1.0 g of 5 % palladium on carbon, and 100 ml of dichloromethane is hydrogenated in a Parr apparatus (50 psi, ambient temperature). The catalyst is then filtered off, and the filtrate is diluted with heptane (50 ml). The dichloromethane-heptane solution is fractionally distilled (bath temperature 50°C) until the product begins to crystallize, then cooled and filtered to give 11.9 g (58 % yield) of the title compound, as a white crystalline solid, m.p. 153-155°C.

Anal. Calcd. for $C_{22}H_{40}N_2O_6$: C, 61.7; H, 9.4; N, 6.5.

Found: C, 61.5; H, 9.2; N, 6.6.

The compounds in the following examples are prepared according to the procedure of Example 18 utilizing the appropriate starting materials.

Example 19: Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) n-butylmalonate
m.p. 92-93°C

Anal. Calcd. for $C_{25}H_{46}N_2O_6$: C, 63.8; H, 9.9; N, 5.9;

Found: C, 63.9; H, 9.8; N, 6.0.

Example 20: Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) diethylmalonate
m.p. 115-118°C

Anal. Calcd. for $C_{25}H_{46}N_2O_6$: C, 63.8; H, 9.9; N, 5.9;

Found: C, 63.9; H, 10.0; N, 5.9.

Example 21: 1-Hydroxy-2,2,6,6-tetramethylpiperidin-4-yl
4-(4-hydroxy-3,5-di-tert-butylbenzoyloxy)-3,5-di-tert-butylbenzoate
m.p. 170-172°C

Anal. Calcd. for $C_{39}H_{59}NO_6$: C, 73.5; H, 9.3; N, 2.2;

Found: C, 72.7; H, 9.2; N, 2.3.

Example 22 : 1-Hydroxy-2,2,6,6-tetramethylpiperidin-4-yl
[4-(2-oxoazepin-1-yl)-2,2,6,6-tetramethylpiperidin-1-yl]acetate
m.p. 251-252°C

Anal. Calcd. for $C_{26}H_{47}N_3O_4$: C, 67.1; H, 10.2; N, 9.0;

Found: C, 67.1; H, 9.9; N, 8.9.

Example 23: Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl)
2-(4-hydroxy-3,5-di-tert-butylbenzyl)-2-n-butylmalonate
m.p. 148-152°C

Anal. Calcd. for $C_{40}H_{68}N_2O_7$: C, 69.7; H, 9.9; N, 4.1;

Found: C, 69.9; H, 10.0; N, 4.2.

Example 24: Tetrakis(1-hydroxy-2,2,6,6-tetramethyl-4-piperidyl) 1,2,3,4-butane-tetracarboxylate
m.p. 201-204°C

Anal. Calcd. for $C_{44}H_{78}N_4O_{12}$: C, 61.8; H, 9.2; N, 6.5

Found: C, 61.5; H, 9.2; N, 6.4.

Example 25: Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) trans-1,4-cyclohexane-dicarboxylate
m.p. 185-192°C

Anal. Calcd. for $C_{26}H_{46}N_2O_6$: C, 64.7; H, 9.6; N, 5.8;

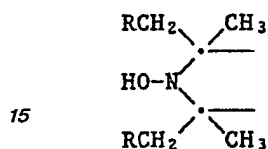
Found: C, 64.3; H, 9.4; N, 5.6.

Claims

5

1. A stabilized ambient curable or acid catalyzed thermosetting coating composition containing an effective stabilizing amount of a hydroxyl substituted hindered amine compound containing the group

10



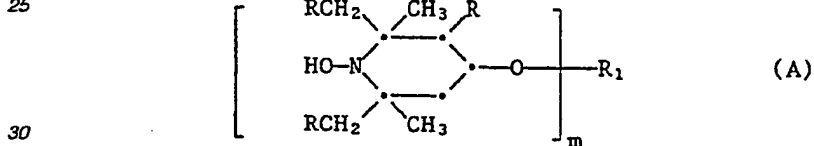
15

where R is hydrogen or methyl.

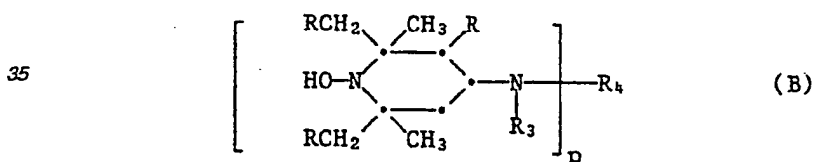
20

2. A composition of claim 1, which contains hindered amine derivative corresponding to one of the formulae A-M

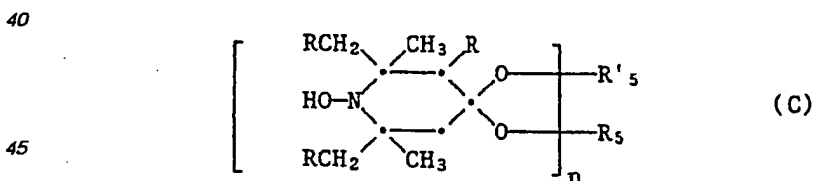
25



30

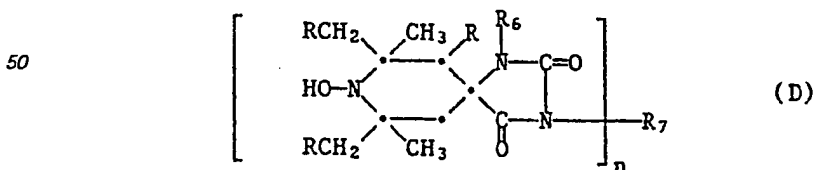


35



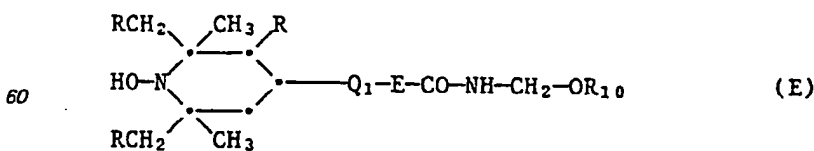
40

45



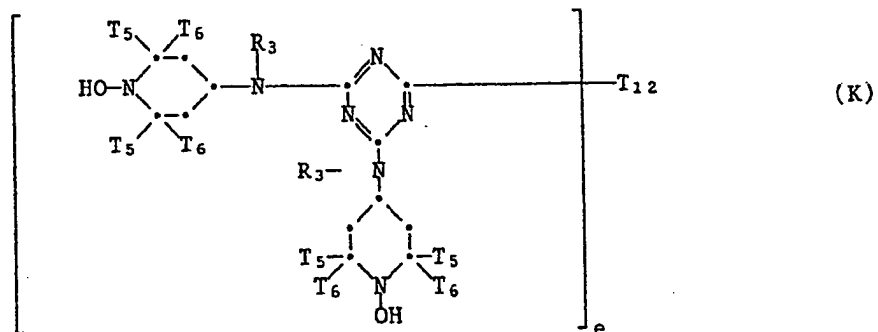
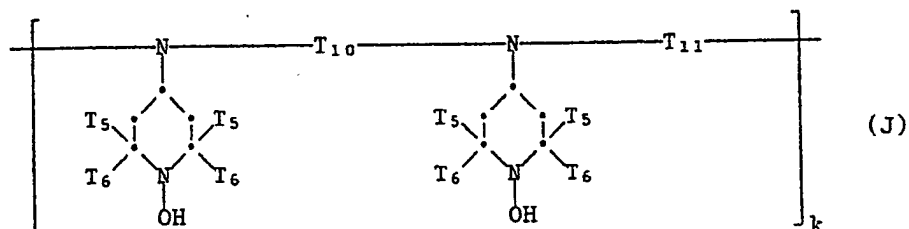
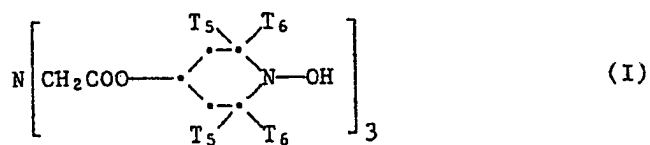
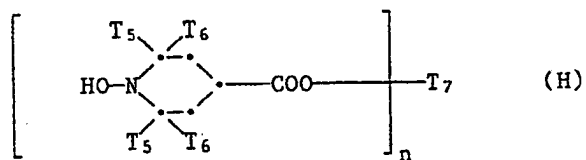
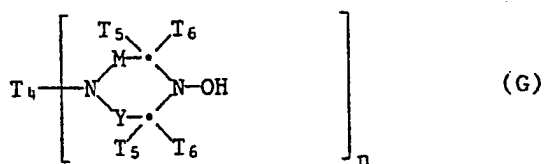
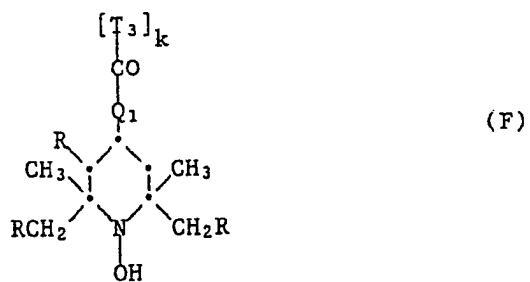
50

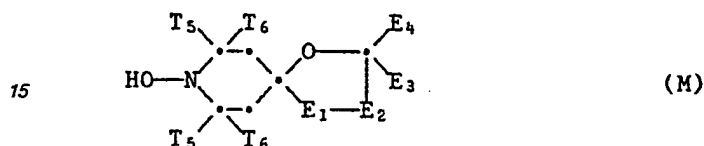
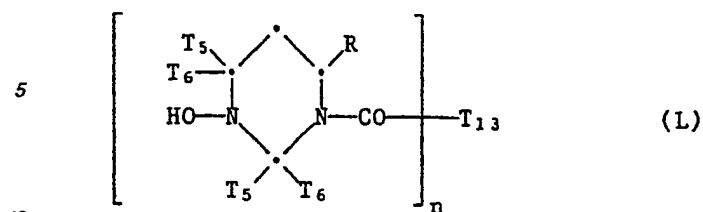
55



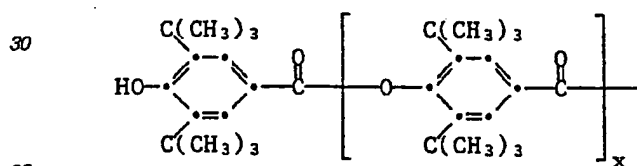
60

65

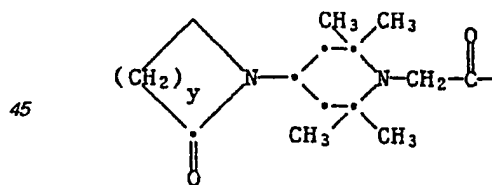




20 wherein
R is hydrogen or methyl,
m is 1-4,
when m is 1,
R₁ is hydrogen, C₁-C₁₈ alkyl optionally interrupted by one or more oxygen atoms, C₂-C₁₂ alkenyl, C₆-C₁₀
25 aryl, C₇-C₁₂ aralkyl, glycidyl, a monovalent acyl radical of an aliphatic, cycloaliphatic, araliphatic or
aromatic carboxylic or carbamic acid; or R₁ is a group



wherein x is 0 or 1, or is a group



50 wherein y is 2-4;
when m is 2,
R₁ is C₁-C₁₂ alkylene, C₄-C₁₂ alkenylene, xylylene, a divalent acyl radical of an aliphatic, cycloaliphatic, araliphatic or aromatic dicarboxylic or dicarbamic acid; or R₁ is a group



wherein D₁ and D₂ are independently hydrogen, C₁-C₈ alkyl, phenyl, benzyl or 3,5-di-tert.butyl-4-hydroxybenzyl and D₃ is an alkyl or alkenyl radical containing up to 18 carbon atoms;
65 when m is 3, R₁ is a trivalent acyl radical of an aliphatic, unsaturated aliphatic, cycloaliphatic, or aromatic

tricarboxylic acid;

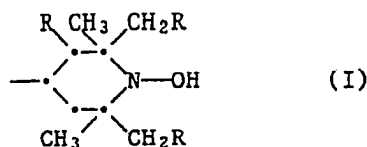
when m is 4, R₁ is a tetravalent acyl radical of a saturated or unsaturated aliphatic or aromatic tetracarboxylic acid;

p is 1, 2 or 3,

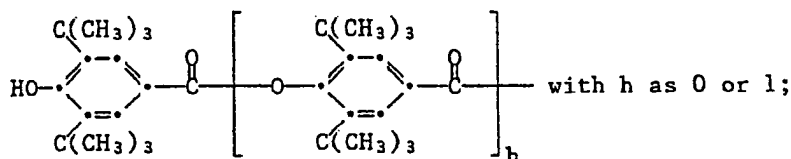
R₃ is hydrogen, C₁-C₁₂ alkyl, C₅-C₇ cycloalkyl, C₇-C₉ aralkyl, C₂-C₁₈ alkanoyl, C₃-C₅ alkenoyl or benzoyl;

when p is 1,

R₄ is hydrogen, C₁-C₁₈ alkyl, C₅-C₇ cycloalkyl, C₂-C₈ alkenyl unsubstituted or substituted by cyano, carbonyl or carbamide group, aryl, aralkyl, or it is glycidyl, a group of the formula -CH₂-CH(OH)-Z or -CONH-Z wherein Z is hydrogen, methyl or phenyl; or R₄ is a group of formula I



or a group of the formula



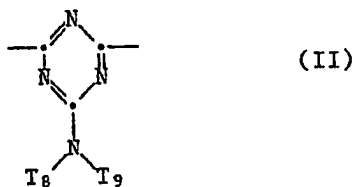
with h as 0 or 1;

or R₃ and R₄ together are alkylene of 4 to 6 carbon atoms or 2-oxo-alkylene or the cyclic acyl radical of an aliphatic or aromatic 1,2- or 1,3-dicarboxylic acid,

when p is 2,

R₄ is C₂-C₁₂ alkylene, C₆-C₁₂ arylene, xylylene, a -CH₂CH(OH)-CH₂- group, or a group -CH₂-CH(OH)-CH₂-O-X-O-CH₂-CH(OH)-CH₂- wherein X is C₂-C₁₀ alkylene, C₆-C₁₅ arylene or C₆-C₁₂ cycloalkylene; or, provided that R₃ is not alkanoyl, alkenoyl or benzoyl, R₄ can also be a divalent acyl radical of an aliphatic, cycloaliphatic or aromatic dicarboxylic acid or dicarbamic acid, or can be the group -CO- or

R₄ is a group of formula II



where T₈ and T₉ are independently hydrogen, alkyl of 1 to 18 carbon atoms or a group of formula I, or T₈ and T₉ together are alkylene of 4 to 6 carbon atoms or 3-oxapentamethylene;

when p is 3,

R₄ is 2,4,6-triazinetriyl,

n is 1 or 2 and

when n is 1,

R₅ and R'₅ are independently C₁-C₁₂ alkyl, C₂-C₁₂ alkenyl, C₇-C₁₂ aralkyl, or R₅ is also hydrogen, or R₅ and R'₅ together are C₂-C₈ alkylene or hydroxyalkylene or C₄-C₂₂ acyloxyalkylene;

when n is 2,

R₅ and R'₅ together are (-CH₂)₂C(CH₂)₂;

R₆ is hydrogen, C₁-C₁₂ alkyl, allyl, benzyl, glycidyl or C₂-C₆ alkoxyalkyl;

when n is 1,

R₇ is hydrogen, C₁-C₁₂ alkyl, C₃-C₅ alkenyl, C₇-C₉ aralkyl, C₅-C₇ cycloalkyl, C₂-C₄ hydroxyalkyl, C₂-C₆

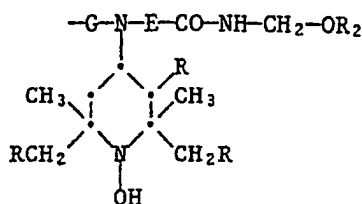
alkoxyalkyl, C₆-C₁₀ aryl, glycidyl, a group of the formula $-(CH_2)_t-COO-Q$ or of the formula $-(CH_2)_t-O-CO-Q$ wherein t is 1 or 2, and Q is C₁-C₄ alkyl or phenyl; or when n is 2,

R₇ is C₂-C₁₂ alkylene, C₆-C₁₂ arylene, a group $-CH_2CH(OH)-CH_2-O-X-O-CH_2-CH(OH)-CH_2-$ wherein X is C₂-C₁₀ alkylene, C₆-C₁₅ arylene or C₆-C₁₂ cycloalkylene, or a group $-CH_2CH(OZ')CH_2-(OCH_2-CH(OZ')CH_2)_2-$ wherein Z' is hydrogen, C₁-C₁₈ alkyl, allyl, benzyl, C₂-C₁₂ alkanoyl or benzoyl; Q₁ is -N(R₈)- or -O-;

E is C₁-C₃ alkylene, the group $-CH_2-CH(R_9)-O-$ wherein R₉ is hydrogen, methyl or phenyl, the group $-(CH_2)_3-NH-$ or a direct bond;

R₁₀ is hydrogen or C₁-C₁₈ alkyl,

R₈ is hydrogen, C₁-C₁₈ alkyl, C₅-C₇ cycloalkyl, C₇-C₁₂ aralkyl, cyanoethyl, C₆-C₁₀ aryl, the group $-CH_2-CH(R_9)-OH$ wherein R₉ has the meaning defined above; or R₈ is a group of the formula I or a group of the formula



wherein G is C₂-C₆ alkylene or C₆-C₁₂ arylene, or R₈ is a group $-E-CO-NH-CH_2-OR_{10}$;

T₃ is ethylene or 1,2-propylene, or is the repeating structural unit derived from an alpha-olefin copolymer with an alkyl acrylate or methacrylate;

k is 2 to 100;

T₄ has the same meaning as R₄ when p is 1 or 2,

T₅ is methyl,

T₆ is methyl or ethyl, or T₅ and T₆ together are tetramethylene or pentamethylene;

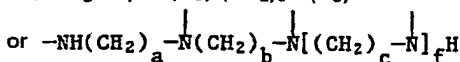
M and Y are independently methylene or carbonyl;

T₇ is the same as R₇;

T₁₀ and T₁₁ are independently alkylene of 2 to 12 carbon atoms, or T₁₁ is a group of formula II;

e is 2, 3 or 4 and

T₁₂ is a group $-N(R_5)-(CH_2)_d-N(R_5)-$



where a, b and c are independently 2 or 3, d is 2 to 10 and f is 0 or 1;

T₁₃ is the same as R₄ with the proviso that T₁₃ cannot be hydrogen when n is 1;

E₁ and E₂, being different, each are $-CO-$ or $-N(E_5)-$, and E₅ is hydrogen, C₁-C₄ alkyl or C₄-C₂₂-alkoxycarbonylalkyl;

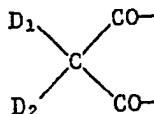
E₃ is hydrogen, alkyl of 1 to 30 carbon atoms, phenyl, naphthyl, said phenyl or said naphthyl substituted by chlorine or by alkyl of 1 to 4 carbon atoms, or phenylalkyl of 7 to 12 carbon atoms, or said phenylalkyl substituted by alkyl of 1 to 4 carbon atoms, and

E₄ is hydrogen, alkyl of 1 to 30 carbon atoms, phenyl, naphthyl or phenylalkyl of 7 to 12 carbon atoms, or

E₃ and E₄ together are polymethylene of 4 to 17 carbon atoms, or said polymethylene substituted by up to four alkyl groups of 1 to 4 carbon atoms.

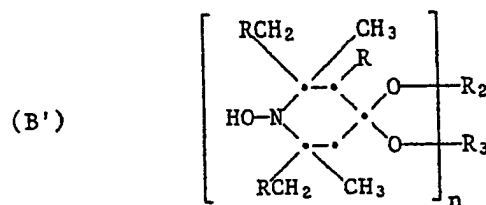
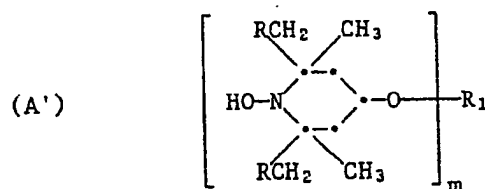
3. A composition according to claim 2, which contains a hindered amine derivative of formula A, B, G-K or M wherein R is hydrogen and T₅ and T₆ are methyl.

4. A composition according to claim 2, which contains a hindered amine derivative of formula A wherein R is hydrogen, m is 1, 2 or 4 and when m is 1, R₁ is an acyl radical of an aliphatic carboxylic acid having 12-18 C atoms or of an aromatic carboxylic acid having 7-15 C atoms and when m is 2, R₁ is a divalent acyl radical of an aliphatic dicarboxylic acid having 4-18 C atoms or of an aromatic dicarboxylic acid having 8-14 C atoms or R₂ is a group



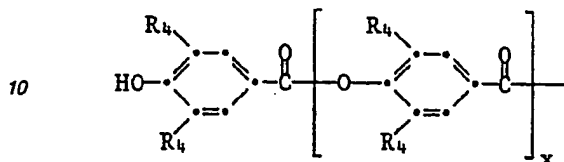
wherein D₁ is C₁-C₈ alkyl or 3,4-di-tert.butyl-4-hydroxybenzyl and D₂ is D₁ or hydrogen, and when m is 4, R₁ is tetravalent acyl radical of a butane- or pentane-tetracarboxylic acid.

5. A composition of claim 1, containing di(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) sebacate.
6. A composition of claim 1, containing 1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl stearate.
7. A composition of claim 1, containing di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) diethylmalonate.
8. A composition of claim 1, containing di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) isophthalate.
9. A composition of claim 1, containing tetrakis (1-hydroxy-2,2,6,6-tetramethyl-4-piperidyl) 1,2,3,4-butane-tetracarboxylate.
10. A composition of claim 1, containing the polycondensation product of 2,4-dichloro-6-tert-octylamino-s-triazine and 4,4'-hexamethylenebis(amino-1-hydroxy-2,2,6,6-tetramethylpiperidine).
11. A composition according to claim 1, wherein the hindered amine derivative is contained in an amount of 0.1 to 10 % by weight, based on resin solids.
12. A composition according to claim 1, which additionally contains a UV absorber selected from the group consisting of benzophenones, benzotriazoles, acrylic acid derivatives, aryl-s-triazines, organic nickel compounds and oxanilides.
13. A composition according to claim 12, which contains a benzotriazole UV absorber selected from the group consisting of 2-[2-hydroxy-3,5-di-(alpha,alpha-dimethylbenzyl)-phenyl]-2H-benzotriazole, 2-(2-hydroxy-3,5-di-tert-octylphenyl)-2H-benzotriazole, 2-(2-hydroxy-3-alpha,alpha-dimethylbenzyl-5-tert-octylphenyl)-2H-benzotriazole, 2-(2-hydroxy-3-tert-octyl-5,alpha,alpha-dimethylbenzylphenyl)-2H-benzotriazole, 2-(2-hydroxy-5-tert-octylphenyl)-2H-benzotriazole, 2-[2-hydroxy-3-tert-butyl-5-(2-(omega-hydroxy-octa(ethyleneoxy)-carbonyl)-ethylphenyl)-2H-benzotriazole, 2-(2-hydroxy-3,5-di-tert-amylphenyl)-2H-benzotriazole, 5-chloro-2-[2-hydroxy-3,5-di(alpha,alpha-dimethylbenzyl)-phenyl]-2H-benzotriazole, 5-chloro-2-(2-hydroxy-3,5-di-tert-butylphenyl)-2H-benzotriazole, 2-[2-hydroxy-3-tert-butyl-5-(2-oxethylcarbonyl)ethylphenyl]-phenyl]-2H-benzotriazole, 2-(2-hydroxy-3-sec.dodecyl-5-methylphenyl)-2H-benzotriazole and hexamethylene di[β-(3-tert.butyl-4-hydroxy-5-[2-benzotriazolyl]-phenyl)-propionate].
14. A coating composition according to claim 1, which is an ambient curable system based on an alkyd resin, thermoplastic acrylic resin, acrylic alkyd resin, polyester resin, or said resins modified with silicones, isocyanates, epoxides, isocyanurates, ketimines or oxazolidines, or the system is based on a cellulose ester or on an epoxide resin.
15. A coating composition according to claim 1, which is an acid catalyzed thermosetting system based on a hot crosslinkable acrylic, polyester, polyurethane, polyamide or alkyd resin.
16. A composition according to claim 1, which is an enamel for industrial finishes.
17. A composition according to claim 1, which is a refinishing enamel for automobiles.
18. A compound corresponding to the formulae A' or B'

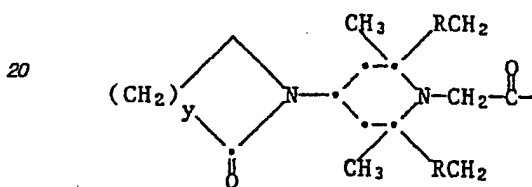


wherein R is hydrogen or methyl;
 m is 1-4;
 when m is 1, R₁ is

5

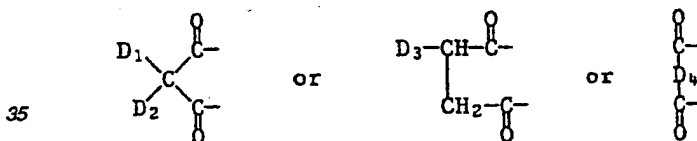


15 wherein x is 0 or 1 and R₄ independently are C₁-C₈alkyl, or



with y is 2-4;
 when m is 2, R₁ is

30



40 wherein D₁ and D₂ are independently C₁-C₈alkyl, C₆-C₁₀aryl, C₇-C₁₂aralkyl or 3,5-di-t-butyl-4-hydroxybenzyl and D₂ is also hydrogen, D₃ is C₁-C₁₈alkyl or C₂-C₁₈alkenyl, and D₄ is C₅-C₁₂cycloalkylene;
 when m is 3, R₁ is a trivalent acyl radical of an aliphatic, unsaturated aliphatic, cycloaliphatic or aromatic tricarboxylic acid having up to 20 C-atoms;
 when m is 4, R₁ is a tetravalent acyl radical of a saturated or unsaturated aliphatic or aromatic tetracarboxylic acid having up to 20 C-atoms;

45 n is 1 or 2;

when n is 1, R₂ and R₃ are independently hydrogen, C₁-C₁₂alkyl, C₂-C₁₂alkenyl or C₇-C₁₂aralkyl; and

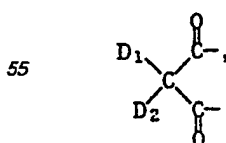
when n is 2, R₂ and R₃ together are (-CH₂)₂C(CH₂)₂.

19. A compound of claim 18, wherein R is hydrogen.

20. A compound of formula A' of claim 18, wherein m is 2,

50

R₁ is



60

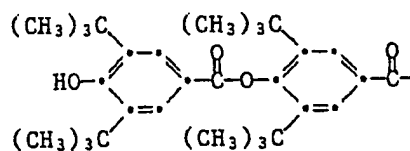
D₁ is C₁-C₈alkyl or 3,5-di-t-butyl-4-hydroxybenzyl and
 D₂ is D₁ or hydrogen.

21. A compound of formula A' of claim 18, wherein

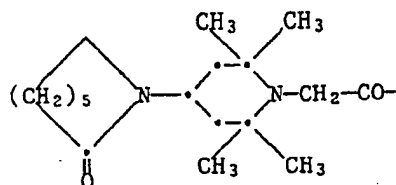
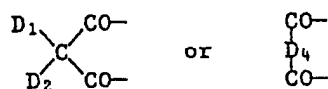
R is hydrogen, m is 1, 2 or 4 and

65

when m is 1, R₁ is a group



or

when m is 2, R₁ is a group

wherein D₁ is C₁-C₄alkyl or 3,5-di-*t*-butyl-4-hydroxybenzyl and D₂ is D₁ or hydrogen and D₄ is cyclohexylene,

and when m is 4, R₁ is the tetravalent acyl radical of a butanetetracarboxylic acid.

22. Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) methylmalonate according to claim 18.

23. Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) *n*-butylmalonate according to claim 18.

24. Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) diethylmalonate according to claim 18.

25. Di-(1-hydroxy-2,2,6,6-tetramethylpiperidin-4-yl) 2-(4-hydroxy-3,5-di-*tert*-butylbenzyl)-2-*n*-butylmalonate according to claim 18.

26. Tetrakis(1-hydroxy-2,2,6,6-tetramethyl-4-piperidyl) 1,2,3,4-butanetetracarboxylate according to claim 18.

27. Use of a compound of claim 18 as stabilizer for organic materials against actinic degradation.

28. Use according to claim 27 as stabilizer for organic polymers.

29. Use according to claim 27 as stabilizer for photographic layers.



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 88 81 0622

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 4)
A	EP-A-0 157 738 (CIBA-GEIGY AG) * Claims; page 6, lines 9-11; example 7 *	1,18	C 07 D 401/14 C 07 D 401/12 C 07 D 211/94
A	EP-A-0 082 244 (AMERICAN CYANAMID CO.) * Page 4, lines 21-23; page 7, lines 4-27; claims * -----	1	C 09 D 7/12 C 08 K 5/34 C 08 K 5/35
			TECHNICAL FIELDS SEARCHED (Int. Cl. 4)
			C 07 D C 09 D C 08 K C 08 L
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 12-12-1988	Examiner DE LOS ARCOS Y VELAZQUEZ
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

EPO FORM 1503 03.82 (P0401)